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WORLD INTELLECTUAL PROPERTY ORGANIZATION-International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:
 A01N 37/18, 43/04, C12Q 1/00, 1/02, 1/68, C12N 5/00, 5/06, 15/00, 15/06, 15/09, 15/10, 15/11, G01N 33/53

(11) International Publication Number:

WO 98/54963

A2 |

(43) International Publication Date:

10 December 1998 (10.12.98)

(21) International Application Number:

PCT/US98/11422

(22) International Filing Date:

4 June 1998 (04.06.98)

(30) Priority Data:

60/048,915 60/048,882 6 June 1997 (06.06.97) 6 June 1997 (06.06.97) US

(Continued on the following page)

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- (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TI, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).

Published

With declaration under Article 17(2)(a); without abstract; title not checked by the International Searching Authority.

(54) Title: 207 HUMAN SECRETED PROTEINS

(Continued)

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FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

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207 Human Secreted Proteins

Field of the Invention

This invention relates to newly identified polynucleotides and the polypeptides encoded by these polynucleotides, uses of such polynucleotides and polypeptides, and their production.

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Background of the Invention

Unlike bacterium, which exist as a single compartment surrounded by a membrane, human cells and other eucaryotes are subdivided by membranes into many functionally distinct compartments. Each membrane-bounded compartment, or organelle, contains different proteins essential for the function of the organelle. The cell uses "sorting signals," which are amino acid motifs located within the protein, to target proteins to particular cellular organelles.

One type of sorting signal, called a signal sequence, a signal peptide, or a leader sequence, directs a class of proteins to an organelle called the endoplasmic reticulum (ER). The ER separates the membrane-bounded proteins from all other types of proteins. Once localized to the ER, both groups of proteins can be further directed to another organelle called the Golgi apparatus. Here, the Golgi distributes the proteins to vesicles, including secretory vesicles, the cell membrane, lysosomes, and the other organelles.

Proteins targeted to the ER by a signal sequence can be released into the extracellular space as a secreted protein. For example, vesicles containing secreted proteins can fuse with the cell membrane and release their contents into the extracellular space - a process called exocytosis. Exocytosis can occur constitutively or after receipt of a triggering signal. In the latter case, the proteins are stored in secretory vesicles (or secretory granules) until exocytosis is triggered. Similarly, proteins residing on the cell membrane can also be secreted into the extracellular space by proteolytic cleavage of a "linker" holding the protein to the membrane.

Despite the great progress made in recent years, only a small number of genes encoding human secreted proteins have been identified. These secreted proteins include the commercially valuable human insulin, interferon, Factor VIII, human growth hormone, tissue plasminogen activator, and erythropoeitin. Thus, in light of the pervasive role of secreted proteins in human physiology, a need exists for identifying and characterizing novel human secreted proteins and the genes that encode them. This knowledge will allow one to detect, to treat, and to prevent medical disorders by using secreted proteins or the genes that encode them.

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Summary of the Invention

The present invention relates to novel polynucleotides and the encoded polypeptides. Moreover, the present invention relates to vectors, host cells, antibodies, and recombinant methods for producing the polypeptides and polynucleotides. Also provided are diagnostic methods for detecting disorders related to the polypeptides, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying binding partners of the polypeptides.

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Detailed Description

Definitions

The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

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In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

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As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence contained in SEQ ID NO:X or the cDNA contained within the clone deposited with the ATCC. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the translated amino acid sequence generated from the polynucleotide as broadly defined.

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In the present invention, the full length sequence identified as SEQ ID NO:X was often generated by overlapping sequences contained in multiple clones (contig

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analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X was deposited with the American Type Culture Collection ("ATCC"). As shown in Table 1, each clone is identified by a cDNA Clone ID (Identifier) and the ATCC Deposit Number. The ATCC is located at 10801 University Boulevard, Manassas, Virginia 20110-2209, USA. The ATCC deposit was made pursuant to the terms of the Budapest Treaty on the international recognition of the deposit of microorganisms for purposes of patent procedure.

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A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, the complement thereof, or the cDNA within the clone deposited with the ATCC. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 μg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH₂PO₄; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a

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complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

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The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single-and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine,

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formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990); Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

"SEQ ID NO:X" refers to a polynucleotide sequence while "SEQ ID NO:Y" refers to a polypeptide sequence, both sequences identified by an integer specified in Table 1.

"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.)

25 Polynucleotides and Polypeptides of the Invention

FEATURES OF PROTEIN ENCODED BY GENE NO: 1

This gene is expressed primarily in melanocytes and, to a lesser extent, in testes, ovary, kidney and other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer, disorders of neural crest derived cells including pigmentation defects, melanoma, reproductive organ defects, and defects of the kidney. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skin,

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reproductive, and renal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating disorders that arise from alterations in the number or fate of neural crest derived cells including cancers such as melanoma and defects of the developing reproductive system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 2

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This gene is expressed primarily in infant brain and fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental disorders of the brain or lung. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous and pulmonary systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating or diagnosing disorders associated with abnormal proliferation of cells in the Central nervous system and developing lung.

FEATURES OF PROTEIN ENCODED BY GENE NO: 3

This gene is expressed primarily in breast lymph node and to a lesser extent in ovarian cancer and chondrosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune responses such as inflammation or immune surveillance for

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tumors. This gene may be important for inflammatory responses associated with tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 236 as residues: Lys-45 to Val-50, Lys-69 to Arg-76.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment or diagnosis of immune responses including those associated with tumor-induced inflammation.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 4

This gene is expressed primarily in T-cells and T-cell lymphomas.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunilogical diseases involving T-cells such as inflammation, autoimmunity, and cancers including T-cell lymphomas. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of T-cells and other cells of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing and treating T-cell based disorders such as inflammatory diseases, autoimmune disease and tumors including T-cell lymphomas.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 5

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This gene is expressed primarily in activated monocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation, autoimmunity, infection, or disorders involving activation of monocytes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 238 as residues: Asp-19 to Arg-31.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing or treating diseases that result in activation of monocytes including infections, inflammatory responses or autoimmune diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 6

The translation product of this gene shares sequence homology with terminal deoxynucleotidyltransferase which is thought to be important in catalyzing the elongation of oligo- or polydeoxynucleotide chains.

This gene is expressed primarily in activated human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer, particularly those of the blood such as leukemia and deficiencies in neutrophils such as neutropenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to terminal deoxynucleotidyltransferase indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and differential diagnosis of acute leukemia's. Alternatively, this gene may function in the proliferation of neutrophils and be useful as a treatment for neutropenia, for example, following neutropenia as a result of chemotherapy.

FEATURES OF PROTEIN ENCODED BY GENE NO: 7

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The contig exhibits a reasonable homology to the human chorionic gonadotropic (HCG) analogue-GT beta-subunit as disclosed in U.S. Patent No. 5,508,261 and PCT Publication No. WO 92/22568. There is a high degree of conservation of the structurally important cysteine residues in these identities.

This gene is expressed primarily in IL-1 and LPS induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

FEATURES OF PROTEIN ENCODED BY GENE NO: 8

This gene is expressed primarily in IL-1- and LPS-induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 241 as residues: Ser-14 to Pro-22, Leu-43 to Val-53.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

FEATURES OF PROTEIN ENCODED BY GENE NO: 9

This gene is expressed primarily in IL-1 and LPS induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 242 as residues: Tyr-22 to His-35.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth

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factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

FEATURES OF PROTEIN ENCODED BY GENE NO: 10

This gene is expressed primarily in activated T-cells and to a lesser extent in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune dysfunctions including cancer of the T lymphocytes and autoimmune disorders and inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of immune disorders particularly of T-cell origin and may act as a growth factor for particular subsets of T-cells such as CD4 positive cells which would make this a useful therapeutic for the treatment of HIV and other immune compromising illnesses.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 11

This gene is expressed primarily in fetal tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of many developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing fetus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor or differentiation factor for particular cell types in the developing fetus and may be useful in replacement or other types of therapy in cases where the gene is expressed aberrantly.

FEATURES OF PROTEIN ENCODED BY GENE NO: 12

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This gene is expressed primarily in T-cells and to a lesser extent in tumor tissue including glioblastoma, meningioma, and Wilm's tumor.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system including autoimmune conditions such as rheumatoid arthritis, inflammatory disorders and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 245 as residues: Thr-9 to Ser-14.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis/ modulation of immune function disorders, including rheumatoid arthritis and inflammatory responses.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 13

This gene is expressed primarily in placenta and to a lesser extent in fetal liver and bone marrow.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of hematological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of

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disorders of the above tissues or cells, particularly of the hematological and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematapoietic stem cells or progenitor cells in the treatment of chemotherapy patients or kidney disease.

FEATURES OF PROTEIN ENCODED BY GENE NO: 14

This gene is expressed primarily in stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of hematapoietic disorders including cancer, neutropenia, anemia, and thrombocytopenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematapoietic and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematapoietic stem cells or progenitor cells, in particular following chemotherapy treatment.

FEATURES OF PROTEIN ENCODED BY GENE NO: 15

The translation product of this gene shares sequence homology with epsilon-COP from Bos taurus which is thought to be important as a component of coatomer, a complex of seven proteins, that is the major component of the non-clathrin membrane coat. Preferred polypeptides encoded by this gene comprise the following amino acid sequences:

MAPPAPGPASGGSGEVDELFDVKNAFYIGSYQQCINEAXXVKLSSPERDVERD

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VFLYRAYLAQRKFGVVLDEIKPSSAPELQAVRMFADYLAHESRRDSIVAELDRE MSRSXDVTNTTFLLMAASIYLHDQNPDAALRALHQGDSLECTAMTVQILLKLD RLDLARKELKRMQDLDEDATLTQLATAWVSLATGGEKLQDAYYIFQEMADKCS PTLLLLNGQAACHMAQGRWEAAEGLLQEALDKDSGYPETLVNLIVLSQHLGKP PEVTNRYLSQLKDAHRSHPFIKEYQAKENDFDRLVLQYAPSAEAGPELSGP (SEQ ID NO:458); or RDVERDVFLYRAYLAQRKFGVVLDEIKPSSAPELQAVRMF ADYLAHESRRDSIVAELDREMSRSXDVTNTTFLLMAASIYLHDQNPDAALRALH **QGDSLECTAMTVQILLKLDRLDLARKELKRMQDLDEDATLTQLATAWVSLATG** GEKLQDAYYIFQEMADKCSPTLLLLNGQAACHMAQGRWEAAEGLLQEALDKD SGYPETLVNLIVLSQHLGKPPEVTNRYLSQLKDAHRSHPFIKEYQAKENDFDRL 10 VLQYAPSA (SEQ ID NO:459).

This gene is expressed primarily in activated monocytes and T-cells, and to a lesser extent in multiple other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunomodulation, specifically relating to transport problems in these cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell 20 type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to epsilon-COP indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating /diagnosing problems with the cellular transport of proteins that may result in immunologic dysfunction.

FEATURES OF PROTEIN ENCODED BY GENE NO: 16

The translation product of this gene shares sequence homology with an RNA helicase which is thought to be important in polynucleotide metabolism. The translation product of this contig exhibits good homology to the LbeIF4A antigen of Leishmania braziliensis. The LbeIF4A antigen, or immunogenic portions of it, can be used to induce protective immunity against leishmaniasis, specifically L. donovani, L. chagasi,

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L. infantum, L. major, L. braziliensis, L. panamensis, L. tropica and L. guyanensis. It can also be used diagnostically to detect Leishmania infection or to stimulate a cellular and/or humoral immune response or to stimulate the production of interleukin-12.

This gene is expressed primarily in colon cancer and to a lesser extent in pituitary.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of cancers particularly of the colon. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 249 as residues: Glu-93 to Ala-98, Gln-150 to Leu-156, Leu-220 to Leu-231, Leu-268 to Arg-273, Val-324 to Pro-341, Arg-372 to Asn-380, Ser-405 to Gly-410, Phe-426 to Ala-433, Glu-458 to Asp-470, Arg-506 to Ser-547.

The tissue distribution and homology to RNA helicase indicates that polynucleotides and polypeptides corresponding to this gene are useful for development of diagnostic tests for colon cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 17

The translation product of this contig has sequence homology to a cytoplasmic protein that binds specifically to JNK designated the JNK interacting protein-1 or JIP-1 in mice. JIP-1 caused cytoplasmic retention of JNK and inhibition of JNK-regulated gene expression.

This gene is expressed primarily in brain including pituitary cerebellum frontal cortex, fetal brain and to a lesser extent in the kidney cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of the central nervous system disorders including ischemia, epilepsy, Parkinson's disease, and schizophrenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological

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probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Furthermore, the translation product of this contig may suppress the effects of the JNK signaling pathway on cellular proliferation, including transformation by the Bcr-Abl oncogene. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 250 as residues: Pro-6 to Ser-26, Ala-30 to Asp-41, Gly-55 to Ser-61, Gly-74 to Thr-80, Tyr-117 to Ala-123, Tyr-167 to Asp-172, Ala-212 to Cys-223, Pro-239 to Tyr-244.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for enhanced survival and/or differentiation of neurons as a treatment for neurodegenerative disease.

FEATURES OF PROTEIN ENCODED BY GENE NO: 18

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The translation product of this gene shares sequence homology with a liver stage antigen from a protozoan parasite.

This gene is expressed primarily in fetal tissue and to a lesser extent in activated T-cells and other immune cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities and diseases of immune function. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a protozoan antigen indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/immune modulation of parasitic infections.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 19

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Preferred polypeptide encoded by this gene comprise the following polypeptide sequences:

MKAIGIEPSLATYHHIIRLFDQPGDPLKRSSFIIYDIMNELMGKRFSPKD
PDDDKFFQSAMSICSSLRDLELAYQVHGLLKTGDNWKFIGPDQHRNFYYSKFF
DLICLMEQIDVTLKWYEDLIPSAYFPHSQTMIHLLQALDVANRLEVIPKIWER
(SEQ ID NO:460); and/or KDSKEYGHTFRSDLREEILMLMARDKHPPELQVAF
ADCAADIKSAYESQPIRQTAQDWPATSLNCIAILFLRAGRTQEAWKMLGLFRKH
NKIPRSELLNELMDSAKVSNSPSQAIEVVELASAFSLPICEGLTQRVMSDFAINQ
EQKEALSNLTALTSDSDTDSSSDSDSDTSEGK (SEQ ID NO:461). Polynucleotides
encoding such polypeptides are also provided.

This gene is expressed primarily in stromal and CD34 depleted bone marrow cells and to a lesser extent in tissues of embryonic origin.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of hematologic origin including cancers and immune dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematapoietic and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 252 as residues: Ser-28 to Gln-34.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematopoietic stem cells or progenitor cells which may be useful in the treatment of chemotherapy patients suffering from neutropenia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 20

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Gln-26 to Lys-34.

Preferred polypeptide fragments can be found in an alternative open reading frame. These preferred polypeptides comprise the amino acid sequence: MSSDNESDIEDEDLKLELRRLRDKHLKEIQDLQSRQKHEIESLYTKLGKVPPAVI IPPAAPLSGRRRPTKSKGSKSSRSSSLGNKSPQLSGNLSGQSAASVLHPQOTL HPPGNIPESGQNQLLQPLKPSPSSDNLYSAFTSDGAISVPSLSAPGQGTSSTNTV GATVNSQAAQAQPPAMTSSRKGTFTDDLHKLVDNWARDAMNLSGRRGSKGH MNYEGPGMARKFSAPGQLCISMTSNLGGSAPISAASATSLGHFTKSMCPPOOY GFPATPFGAQWSGTGGPAPQPLGQFQPVGTASLQNFNISNLQKSISNPPGSNL RTT (SEQ ID NO:462); IQDLQSRQKHEIESLYTKLGKVPPAVIIPPAAPLSGRRRR PTKSKGSKSSRSSSLGNKSPQLSGNLSGQSAASVLHPQQTLHPPGNIPESGQN QLLQPLKPSPSSDNLYSAFTSDGAISVPSLSAPGQGTSST (SEQ ID NO:463); TSDGAISVPSLSAPGQGTSSTNTVGATVNSQAAQAQPPAMTSSRKGTFTDDLH (SEQ ID NO:464); KGHMNYEGPGMARKFSAPGQLCISMTSNLGGSAPISAAS ATSLGHFTK (SEQ ID NO:465); QPLKPSPSSDNLYSAFTSDGAISVPSLSAPG (SEQ ID NO:466). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in fetal liver and tissues associated with the CNS. Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver and CNS diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver and CNS, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 253 as residues:

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for liver diseases such as hepatocellular carcinomas and diseases of the CNS.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 21

In an alternative reading frame, this gene shows sequence homology to two recently cloned genes, karyopherin beta 3 and Ran_GTP binding protein 5. (See Accession Nos. gil2102696 and gnllPIDle328731.) The Ran_GTP binding protein is related to importin-beta, the key mediator of nuclear localization signal (NLS)-dependent nuclear transport. Based on homology, it is likely that this gene may activity similar to the RAN_GTP binding protein. Preferred polypeptide fragments comprise the amino acid sequence: VRVAAAESMXLLLECAXVRGPEYLTQMWHFMCDALIKA IGTEPDSDVLSEIMHSFAK (SEQ ID NO:467). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in thymus tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for immune disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 22

This gene is expressed primarily in prostate and osteoclastoma tissues. Preferred polypeptide fragments also comprise the amino acid sequence:

MEINNQNCFIVIDLVRTVMENGVEGLLIFGAFLPESWLIGVRCSSEPPKALLLIL

AHSQKRRLDGWSFIRHLRVHYCVSLTIHFS (SEQ ID NO:468). Also preferred are polynucleotide sequences encoding this polypeptide fragment.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, bone and prostate diseases, and cancers, particularly of the bone and prostate. Similarly, polypeptides and antibodies directed to these polypeptides are

useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the bone and prostate systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 255 as residues: Met-1 to Ser-11.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for bone and prostate disorders, especially cancers of those systems.

FEATURES OF PROTEIN ENCODED BY GENE NO: 23

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This gene shares sequence homology with the FK506-binding protein (FKBP-13) family, a known cytosolic receptor for the immunosuppressants. Recently, another group has cloned a very similar gene, recognizing the homology to FK506-binding protein family, calling their gene FKBP23. (See Accession No. 2827255.)

This gene is expressed primarily in lymphoid tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample, especially for those susceptible to immune suppressant therapies and for diagnosis of diseases and conditions, which include, but are not limited to, immune suppressant disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 256 as residues: Ala-19 to Val-31, Arg-38 to Gly-49, Ala-61 to Lys-66, Tyr-68 to Pro-78, Gly-116 to Ala-121, Asp-154 to Ser-162, Glu-173 to Gln-186, Phe-194 to Gly-203, Pro-207 to Val-212.

The tissue distribution and homology to FKBP-12 and -13 indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for immune suppressant disorders.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 24

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This gene is expressed primarily in the brain and in the retina. This gene maps to chromosome 8, and therefore can be used in linkage analysis as a marker for chromosome 8.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological and ocular associated disease states. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the disorders of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 257 as residues: Cys-34 to Asp-40.

The tissue distribution in retina indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and/or detection of eye disorders including blindness, color blindness, impaired vision, short and long sightedness, retinitis pigmentosa, retinitis proliferans, and retinoblastoma. Expression in the brain indicates a role in the is useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 25

This gene shows sequence homology to a newly identified class of proteins expressed in the nervous system, called stathmin family. (See Accession No. 2585991; see also Eur. J. Biochem. 248 (3), 794-806 (1997).) The stathmin family appears to be an ubiquitous phosphoprotein involved as a relay integrating various intracellular signaling pathways. These pathways affect cell proliferation and differentiation.

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Preferred polypeptide fragments comprise the amino acid sequence:

QDKHAEEVRKNKELKEEASR (SEQ ID NO:469); QQDLSPWAAPVGCPLXXASX

TCHXLPLSGCLRRQSXSLPVVAXLCFWFSCPLASLFVPGQPCVTCPFPSLPFQD

KHAEEVRKNKELKEEASR (SEQ ID NO:470). Also preferred are the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 26

The polynucleotide sequence of this gene contains a domain similar to a Flt3 ligand peptide. Preferred polypeptide fragments comprise the amino acid sequence: PTRCCTTQPCRSSARRPCWVPMVPSPEGREXQPTCPS (SEQ ID NO:471). Thus, this gene may have activity as binding to Flt3 receptors, a process known to promote angiogenesis and/or lymphangiogenesis.

This gene is expressed in human tonsil, and to a lesser extent in teratocarcinoma, placenta, colon carcinoma, and fetal kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the tonsil, as well as cancers, such as colon, reproductive, and kidney cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful

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in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tonsils, colon, reproductive organs, and kidneys, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 259 as residues: Pro-22 to Glu-33.

The tissue distribution in tonsil and several cancers and fetal tissues indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the tonsil or colon, such as tonsillitis, inflammatory diseases involving nose and paranasal sinuses, especially during the infection of influenza, adenoviruses, parainfluenza, rhinoviruses. The gene may also be useful in the diagnosis and treatment of neoplasms of nasopharynx or colon origins.

FEATURES OF PROTEIN ENCODED BY GENE NO: 27

In an alternative reading frame exists a large open reading frame that encodes a preferred polypeptide. Preferred polypeptide fragments comprise the amino acid sequence:

MKRSLNENSARSTAGCLPVPLFNQKKRNRQPLTSNPLKDDSGISTPSDNYDFP PLPTDWAWEAVNPEXAPVMKTVDTGQIPHSVSRPLRSQDSVFNSIQSNTGRSQ GGWSYRDGNKNTSLKTWXKNDFKPQCKRTNLVANDGKNSCPMSSGAQQQK QLRTPEPPNLSRNKETELLRQTHSSKISGCTMRGLDKNSALQTLKPNFQQNQY KXQMLDDIPEDNTLKETSLYQLQFKEKASSLRIISAVIESMKYWREHAQKTVLL FEVLAVLDSAVTPGPYYSKTFLMRDGKNTLPCVFYEIDRELPRLIRGRVHRCVG NYDQKKNIFQCVSVRPASVSEQKTFQAFVKIADVEMQYYINVMNET (SEQ ID NO:472); SQDSVFNSIQSNTGRSQGGWSYRDGNKNTSLKTWXKNDFKPQCKR (SEQ ID NO:473); NKETELLRQTHSSKISGCTMRGLDKNSALQTLKPNF (SEQ ID NO:474);SSLRIISAVIESMKYWREHAQKTVLLFEVLAVLDSAVTPGPYYSKTFLM (SEQ ID NO:475); and PRLIRGRVHRCVGNYDQKKNIFQCVSVRPASVSEQKT FQAFV (SEQ ID NO:476).

This gene is expressed primarily in human testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, male reproductive disorders, including cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a hormone with reproductive or other systemic functions; contraceptive development; male infertility of testicular causes, such as Kleinfelterís syndrome, varicocele, orchitis; male sexual dysfunctions; testicular neoplasms; and inflammatory disorders such as epididymitis.

FEATURES OF PROTEIN ENCODED BY GENE NO: 28

This gene is expressed primarily in apoptotic T-cell.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases relating to T cells, as well as cancer in general. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the disorders of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune disorders. Moreover, since the gene was isolated from an apoptotic cell and based on the understanding of the relationship of apoptosis and cancer, it is likely that this gene may play a role in the genesis of cancer.

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polypeptide fragments. This gene maps to human chromosome 11, and therefore is useful in linkage analysis as a marker for chromosome 11.

This gene is expressed primarily in human T cells and to a lesser extent in human colon carcinoma.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 263 as residues: Leu-21 to Ala-30, Ser-38 to Asp-47, Pro-87 to Asp-94, Leu-197 to Thr-204, Pro-256 to Ser-262, Thr-277 to Arg-282, Thr-293 to Trp-303.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immune disorders and gastrointestinal diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 31

The translation product of this gene shares sequence homology with Ribosomal protein L11 of Caenorhabditis elegans. (See Accession No. 156201.) Preferred polypeptide fragments comprise the amino acid sequence:

ERGVSINQFCKEFNERTKDIKEGIPLPTKILVKPDRTFEIKIGQPTVSYFLKAAAG IEKGARQTGKEVAGLVTLKHVYEIARIKAQDEAFALQDVPLSSVVRSIIGSARSL

GIRVVKDLSSEELAAF QKERAIFLAAQKEADLAAQEEAAKK (SEQ ID NO:483). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in human embryo tissue and to a lesser extent in human epithelioid sarcoma and other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development disorders and epithelial cell cancer. Similarly, polypeptides and antibodies

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FEATURES OF PROTEIN ENCODED BY GENE NO: 29

This gene is expressed primarily in human tonsils.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, gastrointestinal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of gastrointestinal diseases.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 30

The translation product of this gene shares sequence homology with C44C1.2 gene product of Caenorhabditis elegans with unknown function. Preferred polypeptide fragments comprise the amino acid sequence:

- GVFRPCVCGRPASLTCSPLDPEVGPYCDTPTMRTLFNLLWLALACSPVHTTLSK
 SDAKKAASKTLLEKSQFSDKPVQDRGLVVTDLKAESVVLEHRSYCSAKARDRH
 FAGDVLGYVTPWNSHGYDVTKVFGSKFTQISPVWLQLKRRGREMFEVTGLHD
 VDQGWMRAVRKHAKGLHIVPRLLFEDWTYDDFRNVLDSEDEIEELSKTVVQVA
 KNQHFDGFVVEVWNQLLSQKRVGLIHMLTHLAEALHQARLLALLVIPPAITPGT
 DQLGMFTHKEFEQLAPVLDGFSLMTYDYSTAHQPGPNAPLSWVRACVQVLDP
 KXKWRTKSSWGSTSMXWTXRXPXDARXPVVGXRXIQXLKDHXPRMVLDSK
- 30 KXKWRTKSSWGSTSMXWTXRXPXDARXPVVGXRXIQXLKDHXPRMVLDSK PQ (SEQ ID NO:477); TCSPLDPEVGPYCDTPTMRTLFNLLWLALACSPVHTTLS (SEQ ID NO:478); LVVTDLKAESVVLEHRSYCSAKARDRHFAGDVLGYVTPW NSHGYDVTKVFGSKF (SEQ ID NO:479); REMFEVTGLHDVDQGWMRAVRK HAKGLHIVPRLLFEDWTYDDFRNVLDSEDE (SEQ ID NO:480); HFDGFVVEVW
- 35 NQLLSQKRVGLIHMLTHLAEALHQARLLALLVIPPAITPGTDQLGM (SEQ ID NO:481); DGFSLMTYDYSTAHQPGPNAPLSWVRACVQVLDPKXKWRTKSSW GST (SEQ ID NO:482). Also preferred are polynucleotide fragments encoding these

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directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the embryonic and epithelial cell systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 264 as residues: Lys-34 to Gly-40.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of developmental disorders and epithelial cancer.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 32

This gene is expressed primarily in resting T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory and general immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of disorders of immune system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 33

This gene is believed to reside on chromosome 1. Accordingly, polynucleotides derived from this gene are useful in linkage analysis as chromosome 1 markers.

This gene is expressed primarily in prostate and to a lesser extent in soares adult brain, human umbilical vein endothelial cells, and amniotic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate-related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the urinary system and nervous system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for the diagnosis and treatment of disorders of the urinary and nervous systems.

FEATURES OF PROTEIN ENCODED BY GENE NO: 34

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This gene shares sequence homology with R05G6.4 gene product. (See Accession No. gil1326338.) This gene also shares sequence homology with the cyclophilin-like protein CyP-60. (See Accession No. 1199598, see also Biochem. J. 314 (1), 313-319 (1996).) Preferred polypeptide fragments comprise the amino acid sequence: AVYTYHEKKKDTAASGYGTQNIRLSRDAVKDFDCCCLSLQPCHDPVVTPDGYL YEREAILEYILHQKKEIARQMKAYEKQRGTRREEQKELQRAASQDHVRGFLEKE SAIVSRP LNPFTAKALSGTSPDDVQPGPSVGPPSKDKDKVLPSFWIPSLTPEAK ATKLEKPSRTVTCPMSGKPLRMSDLTPVHFTPLDSSVDRVGLITRSERYVCAVT RDSLSNATPCAVLRPSGAVVTLECVEKLIRKDMVDPVTGDKLTDRDIIVLQRGT (SEQ ID NO:484); YLYEREAILEYILHQKKEIARQMKAYEKQRGTRREEQKELQ RAASQDHVRGFLE (SEQ ID NO:485); and FTAKALSGTSPDDVQPGPSVGPP SKDKDKVLPSFWIPSLTPEAKATKLEKPSRTVTCPMSGKPL (SEQ ID NO:486). Also preferred are polynucleotide fragments that encode these polypeptide fragments.

This gene is expressed primarily in human testis and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders and in particular testicular cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing

immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system. Expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders of the male reproductive system and in particular of testicular cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 35

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The translation product of this gene shares sequence homology with Lpe5p of Saccharomyces cerevisiae which is thought to be important in the metabolism of phospholipids.

This gene is expressed primarily in liver and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, metabolic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and nervous systems expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 268 as residues: Pro-14 to Leu-20, Lys-28 to Asn-38, Arg-109 to Arg-114, Lys-119 to Asn-124, Glu-152 to Leu-157, Pro-172 to Val-180.

The tissue distribution and homology to Lpe5p of Saccharomyces cerevisiae indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of metabolic and nervous disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 36

This gene shares sequence homology with the nuclear ribonucleoprotein U (HNRNP U), encoded by *C. elegans* (See Accession gil1703576.) Preferred polypeptide fragments comprise the amino acid sequence:

5 MDTSENRPENDVPEPPMPIADQVSNDDRPEGSVEDEEKKESSLPKSFKRKISVV
SATKGVPAGNSDTEGGQPGRKRRWGASTATTQKKPSISITTESLKSLIPDIKPL
AGQEAVVDLHADDSRISEDETERNGDDGTHDKGLKICRTVTQVVPAEGQENGQ
REEEEEEKEPEAEPPVPPQVSVEVALPPPAEHEVKKVTLGDTLTRRSISQQKSGV
SITIDDPVRTAQVPSPPRGKISNIVHISNLVRPFTLGQLKELLGRTGTLVEEAFWI
10 DKIKSHCFVTYSTVEEAVATRTALHGVKWPQSNPKFLCADYAEQDELDYHRGL
LVDRPSETKTEEQGIPRPLHPPPPPPVQPPQHPRAEQREQERAVREQWAERERE
MERRERTRSEREWDRDKVREGPRSRSRSRXRRRKERAKSKEKKSEKKEKAQE
EPPAKLLDDLFRKTKAAPCIYWLPLTDSQIVQKEAERAERAKEREKRRKEQEEE
EQKEREKEAERERNRQLEREKRREHSRERDRERERERDRGDRDRDRERDRE
15 RGRERDRRDTKRHSRSRSRSTPVRDRGGR (SEQ ID NO:488). Also preferred are
the polynucleotide fragments encoding this polypeptide fragments.

This gene is expressed primarily in epididymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the male reproductive system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of male reproductive disorders.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 37

This gene is expressed primarily in amygdala.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory diseases and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the amygdala, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of inflammatory diseases and reproductive disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 38

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This gene shares sequence homology with human opsonin protein P35 fragment. (See Accession No. R94181.) The opsonin protein activates the phagocytosis of pathogenic microbes by phagocytic cells. Preferred polypeptide fragments comprise the amino acid sequence: GCDSCPPHLPREAFAQDTQAEGECSSRAERADMCPDAP PSQEVPEGPGAAP (SEQ ID NO:489). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in immune-related tissues such as thymus, macrophage, T cells and to a lesser extent in many other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders and infectious disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and infectious disease, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 271 as residues: Lys-9 to Arg-14, Met-38 to Asp-51.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders, as well as the treatment and/or diagnosis of infectious disease.

FEATURES OF PROTEIN ENCODED BY GENE NO: 39

The translation product of this gene shares sequence homology with alpha-2 type I collagen which is thought to be important in tissue repair. (See, e.g., 211607.) Preferred polypeptide fragments comprise the amino acid sequence: PQLPSCGRPW PGTASVFQSHTQGPREDPDPCRAQGSAGTHCPISLSPPRQ (SEQ ID NO:490). Also preferred are the polynucleotide sequences encoding these polypeptide sequences.

This gene is expressed primarily in the brain and to a lesser extent in the kidney and thymus

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain, kidney, and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, kidney, and immune disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to alpha-2 type I collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tissue repair, and brain, kidney, immune disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 40

The translation product of this gene shares sequence homology with minicollagen which is thought to be important in tissue repair tumor metastasis. (See Accession No. gnllPIDld1006976.) Preferred polypeptide fragments comprise the amino acid sequence: PGFRGPSGSLGCSFFPRSLGRVLPPGCQRPGAHAD 5

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SSPPPTP (SEQ ID NO:491). Also preferred are polynucleotides encoding this polypeptide fragment.

This gene is expressed in ovarian cancer and to a lesser extent in dedritic cells and smooth muscle.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumor metastasis and tissue repair. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumor metastasis and tissue repair, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 273 as residues: Asn-2 to His-11.

The tissue distribution and homology to mini-collegen gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tumor metastasis and tissue repair.

FEATURES OF PROTEIN ENCODED BY GENE NO: 41

This gene shares sequence homology with the HIV TAT protein. (See

25 Accession No. 328416.) Preferred polypeptide fragments comprise the amino acid
sequence: EDLKKPDPASLRAASCGEGKKRKACKNCTCGLAEELEKEK
SREQMSSQPKSACGNCYLGDAFRCASCPYLGMPAFKPGEKVLLS (SEQ ID
NO:492); EDLKKPDPASLRAASCGEGKKRKACKNCTCGLAEELEKEK
SREQMSSQPKSACGNCYLGDAFRCASCPYLGMPAFKPGEKVLLSDSNLHD
30 (SEQ ID NO:493); CGNCYLGDAFRCASCPYLGMPAFKPGEKVLLSDS
(SEQ ID NO:494); SCGEGKKRKACKNCTCGLAEELEKE (SEQ ID NO:495);
SQPKSAC GNCYLGDAFRCASC (SEQ ID NO:496); and REAGQNSERQYVS
LSRD (SEQ ID NO:497). Also preferred are polynucleotide fragments encoding these
polypeptide fragments.

This gene is expressed primarily in the infant brain and to a lesser extent in the breast and testes.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain, testes and breast disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, testes and breast disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 274 as residues: Pro-7 to Val-15.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of brain, testes and breast, and other related disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 42

This gene is expressed primarily in the infant brain, human cerebellum, and to a lesser extent in medulloblastoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain related disorders and medulloblastoma and other brain cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain related disorders and brain cancers, including medulloblastoma, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 275 as residues: Thr-41 to Glu-47.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of human brain related disorders, brain cancers, and medulioblastoma.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 43

The translation product of this gene shares sequence homology with a phosphotyrosine-independent ligand for the lck SH2 domain which is thought to be important in signal transduction related to phosphotyrosine-independent ligand for the lck SH2 domain. (See Accession No. gil1184951.) Preferred polypeptide fragments comprise the amino acid sequence: ESSGQARTLADPGPGWPRQQGMCFGSLT GLSTTPHGFLTVSAEADPRLIESLSQMLSMGFSDEGGWLTRLLQTKNYDIGAAL DTIQYSKH (SEQ ID NO:498). Also preferred are polynucleotide fragments encoding this polypeptide fragment. It is likely that this gene is a new member of a family of phosphotyrosine-independent ligands for the lck SH2 domains.

This gene is expressed primarily in the placenta and to a lesser extent in endothelial cells and neutrophil.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive, cardiovascular, immune, and infectious diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular, reproductive, and immune system, and infectious diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a phosphotyrosine-independent ligand for the lck SH2 domain indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cardiovascular, reproductive, and immune system diseases, as well as infectious diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 44

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This gene is expressed primarily in the fetal brain, cerebellum and to a lesser extent in the placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal cell related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal cell related disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 277 as residues: Thr-20 to Gly-28.

The tissue distribution and homology to proline-rich protein genes indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal cell related disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 45

The translation product of this gene shares sequence homology with precerebellin of human, which is thought to be important in synaptic physiology. (See Accession No. gil180251.) It has been observed that cerebellin-like immunoreactivity is associated with Purkinje cell postsynaptic structures. Thus, it is likely that this gene also have synaptic activity. Preferred polypeptide fragments comprise the amino acid sequence: QEGSEPVLLEGECLVVCEPGRAAAGGPGGAALGEAPPGRVAFXAV RSHHHEPAGETGNGTSGAIYFDQVLVNEGGGFDRASGSFVAPVRGVYSFRFH VVKVYNRQTVQVSLMLNTWPVISAFANDPDVTREAATSSVLLPLDPGDRVSLR LRRGXSTGW (SEQ ID NO:499). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in cerebellum and infant brain. By Northern analysis, a single transcript of 2.4 kb was observed in brain tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, neuronal cell signal transduction and synaptic physiology. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal cell signal transduction and synaptic physiology expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to gene or gene family indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal cell related disorders.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 46

This gene is expressed in fetal liver and spleen, and to a lesser extent in bone marrow, umbilical vein, and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the immune system, particularly hematopoiesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoiesis and immune disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 279 as residues: Asp-30 to Glu-57.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopieotic and immune disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 47

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The translation product of this gene shares sequence homology with a 12 kD nucleic acid binding protein of Feline calcivirus which is thought to be important in viral replication. (See Accession No. 59264)

This gene is expressed primarily in human cardiomyopathy and to a lesser extent in T helper cells, fetal brain and synovial sarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiomyopathy as well as viral infection. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 280 as residues: Trp-20 to Cys-26.

The tissue distribution in cardiomyopathy and homology to viral 12 kD nucleic acid binding protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of cardiomyopathy, including those caused by ischemic, hypertensive, congenital, valvular, or pericardial abnormalities. The gene expression pattern may be the consequence or the cause for these conditions.

FEATURES OF PROTEIN ENCODED BY GENE NO: 48

The translation product of this gene shares sequence homology with tumor necrosis factor related gene product which is thought to be important in tumor necrosis, bacterial and viral infection, immune diseases and immunoreactions.

This gene is expressed primarily in colon and to a lesser extent in ovarian and breast cancers.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of colon, ovary or breast origins. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

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for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the colon, ovary and breast, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Tumor necrosis factors indicates that polynucleotides and polypeptides corresponding to this gene are useful for intervention of cancers of colon, ovary and breast origins, because TNF family members are known to be involved in the tumor development.

FEATURES OF PROTEIN ENCODED BY GENE NO: 49

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The translation product of this gene shares sequence homology with mucins, such as epithelial mucin, which is thought to be important in extracellular matrix functions such as protection, lubrication and cell adhesion (See for example Accession No. R68002). Preferred polypeptide fragments comprise the following amino acid sequence: PRSRPALRPGRQRPPSHSATSGVLRPRKKPDP (SEQ ID NO:500).

20 Also preferred are polynucleotide fragments encoding these polypeptide fragments

Also preferred are polynucleotide fragments encoding these polypeptide fragments. Moreover, this gene maps to chromosome 22q11.2-qter, and therefore, can be used as a marker in linkage analysis for chromosome 22.

This gene is expressed primarily in corpus colosum.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors, especially of corpus colosum, as well as metastatic lesions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the corpus colosum and other solid tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to mucins indicates that polynucleotides and polypeptides corresponding to this gene are useful for serum tumor markers or immunotherapy targets because tumor cells have greatly elevated level of mucin expression and shed the molecules into the epithelial tissues.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 50

This gene is expressed primarily in CD34 depleted buffy coat cord blood and primary dendritic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic disorders and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in CD34 depleted buffy coat cord blood and primary dendritic cells indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopoietic and immune disorders. Secreted or cell surface proteins in the above tissue distribution often are involved in cell activation (e.g. cytokines) or molecules involved in cell surface activation.

FEATURES OF PROTEIN ENCODED BY GENE NO: 51

The translation product of this gene shares sequence homology with Interferon induced 1-8 gene encoded polypeptide which is thought to be important in binding to retroviral rev responsive element. Preferred polypeptide fragment comprise the following amino acid sequences: MTLITPSXKLTFXKGNKSWSSRACSSTLVDP (SEQ ID NO:501). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

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This gene is expressed primarily in CD34 positive cells and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, retroviral infection, such as AIDS, and other immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 284 as residues: Gln-51 to Trp-62.

The tissue distribution and homology to interferon induced gene 1-8 indicates that polynucleotides and polypeptides corresponding to this gene are useful for intervention of retroviral infection including HIV. The factor may be involved in viral stability or viral entry into the cells. Alternatively, the virus/factor complex may elicit the cellular immune reaction.

FEATURES OF PROTEIN ENCODED BY GENE NO: 52

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This gene shares sequence homology to immunoglobulin lambda chain (See Accession No. 2865484). Therefore it is likely that this gene has activity similar to an immunoglobulin lambda chain. Preferred polypeptide fragments comprise the following amino acid sequence: GHPSPALSIAPSDGSQLPCDEVPYGEAHVTRYCKKPLTNS HLETEAQSSSL (SEQ ID NO:502). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in Hodgkin's lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, Hodgkin's lymphoma and other immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 285 as residues: Pro-27 to Thr-32.

The tissue distribution in Hodgkin's lymphoma and the sequence homology indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of Hodgkin's lymphoma, since the elevated expression and secretion by the tumor mass may be indicative of tumors of this type. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 53

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This gene has extensive homology to cDNA for Homo sapiens mRNA for the ISLR gene(See Accession No. AB003184). This protein is considered to be a new member of the Ig superfamily and contains a leucine-rich repeat (LRR) with conserved flanking sequences and a C2-type immunoglobulin (Ig)-like domain. These domains are important for protein-protein interaction or cell adhesion, and therefore it is possible that the novel protein ISLR may also interact with other proteins or cells. The ISLR gene was mapped on human chromosome 15q23-q24 by fluorescence in situ hybridization (See Medline Article No. 97468140). Homology to the ISLR gene has been confirmed by another independent group as well (See Accession No. Hs.102171)

This gene is expressed in a number of tissues including human retina, heart, skeletal muscle, prostate, ovary, small intestine, thyroid, adrenal cortex, testis, stomach, spinal cord, fetal lung and fetal kidney tissues, colon, tonsil and stomach cancer, and to a lesser extent in endometrial stromal cells treated with estradiol, breast tissue, synovium, lymphoma, and number of other tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of colon, ovary and breast origins. However, due to the wide range of expression in various tissues, protein may play a vital role in the development of cancer in other tissues as well, not just those mentioned above. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the colon, ovary and breast, expression of this gene at significantly higher or lower levels may be routinely

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detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Additionally, this gene maps to chromosome 15q23-q24, and therefore, can be used as a marker in linkage analysis for chromosome 15.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 54

This gene is expressed primarily in lung, esophagus, leukemia (Jurkat cells) and breast cancers and to a lesser extent in macrophages treated with GM-CSF fetal tissues and wide range of tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer of wide range of origins. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the solid tumors, lung and leukemia, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Furthermore, due to the high expression level in lung tissue and the proposed function of the multidrug resistance protein 1 gene as the efflux pump responsible for low-drug accumulation in multidrug-resistant cells, protein as well mutants thereof, may also be beneficial as a target for gene therapy, particularly for the chronic patient. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 287 as residues: Met-1 to Lys-16.

The tissue distribution in wide range of cancers and fetal tissues indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection of cells in active proliferation, such as cancers. The gene products may be used for cancer markers or immunotherapy target.

FEATURES OF PROTEIN ENCODED BY GENE NO: 55

This gene maps to the X chromosome.

This gene is expressed primarily in the brain and to a lesser extent in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disease states and developmental disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders, including sex-linked disorders, of the above tissues or cells, particularly of the neurological, developmental systems, and cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Moreover, this gene maps to the X chromosome, and therefore, may be used as a marker in linkage analysis for this chromosome.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, Klinefelter's, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental

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disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 56

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The translation product of this gene shares sequence homology with paxillin which is thought to be important in mediating signal transduction from growth factor receptors to the cytoskeleton. Preferred polynucleotide fragments comprise the following sequence: TGGCTCACTGTCTTACAATCACTGCTGTGGAATCATGA TACCACTTTTAGCTCTTTGCATCTTCCTTCAGTGTATTTTTGTTTTTCAAGAGG GGCTTGTGGTTTCAA (SEQ ID NO:506). Also preferred are polypeptide fragments encoded by these polynucleotide fragments. More preferably, polypeptide fragments comprise the amino acid sequence: LDELMAHLTEMQAKVAVRAD AGKKHLPDKQDHKASLDSMLGGLEQELQDLGIATVPKGHCASCQKPIAGKVI HALGQSWHPEHFVCTHCKEEIGSSPFFERSGLXYCPNDYHQLFSPRCAYCAAP ILDKVLTAMNQTWHPEHFFCSHCGEVFGAEGFHEKDKKPYCRKDFLAMFSPK CGGCNRPVLENYLSAMDTVWHPECFVCGDCFTSFSTGSFFELDGRPFCELHYH HRRGTLCHGCGQPITGRCISAMGYKFHPEHFVCAFCLTQLSKGIFREQNDKTY CQPCFNKLF (SEQ ID NO:507); KASLDSMLGGLEQELQDLGIATVPKGHC ASCQKPIAGKVIHAL (SEQ ID NO:508); CPNDYHQLFSPRCAYCAAPILDKVL TAMNQTWHPEHFFCSHCGEVFGAEG (SEQ ID NO:509); DKKPYCRKDFLAM FSPKCGGCNRPVLENYLSAMDTVWHPECFVCGDCFTSFSTGSFFELDGRPFCE L (SEQ ID NO:510); CGQPITGRCISAMGYKFHPEHFVCAFCLTQLSKGIFRE QNDKTYCQ (SEQ ID NO:511). Polynucleotide fragments encoding these preferred polypeptide fragments are also contemplated.

This gene is expressed primarily in brain, and to a lesser extent in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disease states and developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or

cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Moreover, since this gene shares homology with a gene that maps to chromosome 11, (See Accession No.T87404), gene as well as its translated product may be used for linkage analysis on chromosome 11.

The tissue distribution and homology to paxillin indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and or detection of disease states associated with abnormal signal transduction in brain and/or the developing embryo. This would include treatment or detection of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder and also in the treatment and or detection of embryonic development defects.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 57

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This gene is expressed primarily in fetal spleen, brain, and to a lesser extent in six week old embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders, neurological disorders, and developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and developmental systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 290 as residues: Arg-28 to Gly-34.

The expression of this gene in fetal spleen indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/detection of immune disorders such as arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. In addition the expression of this gene in the early embryo, indicates a key role in embryo development and hence the gene or gene product could be used in the treatment and or detection of embryonic development defects. This would include

treatment or detection of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder and also in the treatment and or detection of embryonic development defects.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 58

The translation product of this gene shares sequence homology with the gene disrupted in the neurodegenerative disease dentatorubal-pallidoluysian atrophy. Moreover a long open reading fame exists in an alternative frame. Preferred polypeptide fragments comprise the following:

MGSSQSVEIPGGGTEGYHVLRVQENSPGHRAGLEPFFDFIVSINGSRLNKDND
TLKDLLKXNVEKPVKMLIYSSKTLELRETSVTPSNLWGGQGLLGVSIRFCSFD
GANENVWHVLEVESNSPAALAGLRPHSDYIIGADTVMNESEDLFSLIETHEAKP
LKLYVYNTDTDNCREVIITPNSAWGGEGSLGCGIGYGYLHRIPTRPFEEGKKIS
LPGQMAGTPITPLKDGFTEVQLSSVNPPSLSPPGTTGIEQSLTGLSISSTPPAVSS
VLSTGVPTVPLLPPQVNQSLTSVPPMNPATTLPGLMPLPAGLPNLPNLNLNLPA
PHIMPGVGLPELVNPGLPPLPSMPPRNLPGIAPLPLPSEFLPSFPLVPESSSAASS
GELLSSLPPTSNAPSDPATTTAKADAASSLTVDVTPPTAKAPTTVEDRVGDSTPV
SEKPVSAAVDANASESP (SEQ ID NO:512); SVEIPGGGTEGYHVLRVQENSPGH
RAGLEPFFDFIVSINGSRLNKDNDTLKDLLKXNVEKPVKMLIYSSKTLELRETS
VTPSNLWGGQGLLGVSIRFCSFDGANENVWH (SEQ ID NO:513); ESNSPAA
LAGLRPHSDYIIGADTVMNESEDLFSLIETHEAKPLKLYVYNTDTDNCREVIITP
NSAWGGEGSLGCGIGYGYLHRIPTRPFEEGKKISLPGQMAGTPITPLKDGFTEV

QLSSVNPPSLSPPGTTGIEQSLTG LSISS (SEQ ID NO:514); RIPTRPFEEGKKI

25 SLPGQMAGTPITPLKDGFTEVQLSSVNPPSLSPPGTTGIEQSLTGLSISSTPPAVS
SVLSTGVPTVPLLPPQVNQSLTSVPPMNPATTLPGLMPLPAGLPNLPNLNLNLP
APHIMPGVGLPELVNPGLPPLPSMPPRN (SEQ ID NO:516); PGLPPLPSMPPRN
LPGIAPLPLPSEFLPSFPLVPESSSAASSGELLSSLPPTSNAPSDPATTTAKADAA
SSLTVDVTPPTAKAPTTVEDRVGDSTPVSEKPVSAAVDAN (SEQ ID NO:517).

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This gene is expressed primarily in prostate cancer, and to a lesser extent in the pineal glands and in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological conditions and pulmonary disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For

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a number of disorders of the above tissues or cells, particularly of the nervous, pulmonary, and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 291 as residues: Asn-9 to Leu-14.

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The abundance of this gene in the pineal gland and its homology to a gene disrupted in the neurodegenerative disease state Dentatorubral-pallidoluysian atrophy indicates that this gene may be useful in the treatment and/or detection of other neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. The abundance of this gene in fetal lung would suggest that misregulation of the expression of this protein product in the adult could lead to lymphoma or sarcoma formation, particularly in the lung; that it may also be involved in predisposition to certain pulmonary defects such as pulmonary edema and embolism, bronchitis and cystic fibrosis; and thus the gen or the gene protein encoded by the gene could be used in the detection and/or treatment of these pulmonary disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 59

This gene is expressed primarily in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developmental system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The expression of this gene primarily in the embryo, indicates the gene plays a key role in embryo development and that the gene or the protein encoded by the gene could be used in the treatment and or detection of developmental defects in the embryo or in infants.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 60

This gene displays homology to nestin, an intermediate filament protein, the expression of which correlates with the proliferation of Central Nervous System progenitor cells and that is useful in the identification of brain tumors. This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. AA527348).

This gene is expressed primarily in kidney and to a lesser extent in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, renal disorders and neurodegenerative conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the excretory and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 293 as residues: Thr-128 to Asn-135.

The tissue distribution and homology to nestin indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and/or treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, its abundance in kidney indicates that it is useful in the treatment and detection of acute renal failure and other disease states associated with the kidney.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 61

Gene shares homology with the latrophilin-related protein 1 precursor as well as the calcium-independent alpha-latrotoxin receptor. Preferred polypeptide fragments

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comprise the following amino acid sequence:

IYKVFRHTAGLKPEVSCFENIRSCARXXXXXXXXXXXXXXXXIFGVLHVVHASVV TAYLFTVSNAFQGMFIFLFLCVLSRKIQEEYYRLFKNVPCC (SEQ ID NO:518); WIFGVLHVVHASVVTAYLFTVSNAFQGMFIFLFLCVLSRKIQEEYYRLFKNVPC C (SEQ ID NO:519). Also preferred are polynucleotide fragments encoding these polypeptide fragments. (See Accession No. 2213659) The translation product of this

gene shares sequence homology with CD 97, a seven transmembrane bound receptor.

This gene is expressed primarily in infant brain and in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders and hematopoeitic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological and hematopoeitic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 294 as residues: Lys-13 to Leu-21.

The tissue distribution of this gene suggest that it may be useful in the detection and/or treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder, while its expression in hematopoietic cell types indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, asthma and immunodeficiency diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 62

This gene is expressed primarily in fetal liver and fetal spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematological and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hematopoetic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 295 as residues: Ser-91 to Lys-98.

The tissue distribution of this gene fetal liver and spleen indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, leukemia, asthma and immunodeficiency diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 63

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Gene shares homology with human serum amyloid protein. Preferred polypeptide fragments comprise the following amino acid sequence:
 ALTRIPPGDWVINVTAVSFAGKTTARFFHSSPPSLGDQARTDPGHQRRD (SEQ ID NO:520) (See Accession No. W13671). Also preferred are polynucleotide fragments encoding these polypeptide fragments This gene maps to chromosome 9, and therefore, may be used as a marker in linkage analysis for chromosome 9 (See Accession No. AA004342).

This gene is expressed primarily in fetal liver and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene in fetal liver-spleen indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, leukemia, asthma, and immunodeficiency diseases.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 64

This gene maps to chromosome 3, and therefore, may be used as a marker in linkage analysis for chromosome 3 (See Accession No. AA219669).

This gene is expressed specifically in the brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disease states. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 65

Gene shares homology with a yeast protein. Preferred polypeptide fragments comprise the following amino acid sequence: LQEVNITLPENSVWYERYKFDIP VFHL (SEQ ID NO:521). Also preferred are polynucleotide fragments encoding these polypeptide fragments. (See Accession No. 1332638)

This gene is expressed primarily in fetal tissue (fetus and fetal liver).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver disorders and cancers (e.g. hepatoblastoma). Similarly,

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polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 298 as residues: Asn-59 to Glu-64.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

FEATURES OF PROTEIN ENCODED BY GENE NO: 66

Gene has homology with a B-cell surface antigen which may indicate gene plays 20 a role in the immune response, including, but not limited to disorders and infections of the immune system. Preferred polynucleotide fragments comprise the following sequence: TAGCATGTAGCCAGTCGAATAACNTATAAGGACAAAGTGGAGTC CACGCGTGCGGCCGTCTAGACTAGTGGATCCCCCGGCTGCAGGATTCGGC 25 ACGAG (SEQ ID NO:523). Also preferred are polypeptide fragments encoded by these polynucleotide fragments (See Accession No.T94535). Additionally, this gene shares homology with an interferon-gamma receptor. Preferred polypeptide fragments also comprise the following amino acid sequence: MQGSGSQFRACLLCLCFSCPC SPGGPRWNSRQGGRRFPKTCRAISQNLVFKYKTFCPVRYMQPHRSSLCLHFTS 30 YVFILSTWGSLRTYSTDLKKKKKNSRGGPVPIRPKS (SEQ ID NO:522); MQGSGSQFRACLLCLCFSCPCSPGGPRWNSRQGGRRFPKTCRAISQNLVFK (SEQ ID NO:524); PVRYMQPHRSSLCLHFTSYVFILSTWGSLRTYSTDLKKKKK NSRGGPVPIRPKS (SEQ ID NO:525); and GEEQRDCSLGWRGVGMRATHCQAA RMFVLFSLPKYAGL (SEQ ID NO:526). Also preferred are polynucleotide fragments 35 encoding these polypeptide fragments

This gene is expressed primarily in T-cells and gall bladder.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological disorders and conditions (immunodeficiencies, cancer, leukemia, hematopoeisis). Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and digestive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 299 as residues: Thr-41 to Gly-52.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of immune disorders including: leukemias, lymphomas, auto-immune disorders, immuno-supressive (transplantation) and immunodeficiencies (e.g. AIDS), inflammation and hematopoeitic disorders. The expression of this gene in gall bladder would suggest a possible role for this gene product in digestive disorders, particularly of the pancreas.

FEATURES OF PROTEIN ENCODED BY GENE NO: 67

This gene maps to chromosome 11, and therefore, may be used as a marker in linkage analysis for chromosome 11 (See Accession No. AA011622).

This gene is expressed primarily in a variety of fetal and developmental tissues (e.g. fetal spleen, infant brain).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental, immune or neurological abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing immune and central nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 300 as residues: Ser-38 to Ser-43.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for developmental abnormalities or fetal deficiencies. The detection in infant brain would suggest a role in neurological disorders (both developmental and neurodegenerative conditions of the brain and nervous system, behavioral disorders, depression, schizophrenia, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia). In addition, the detection in spleen would similarly suggest a role in detection and treatment of immunologically mediated disorders (e.g. immunodeficiency, inflammation, cancer, wound healing, tissue repair, hematopoeisis).

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 68

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This gene is expressed primarily in spleen, T-cells, and fetal heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological deficiencies, including AIDSand cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and cardiovascular systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immune disorders including: leukemias, lymphomas, autoimmune disorders, immunodeficiencies (e.g. AIDS), immuno-suppressive conditions (transplantation) and hematopoeitic disorders. The expression in fetal heart indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cadiovascular disorders (e.g. heart disease, restenosis, atherosclerosis, stoke, angina, thrombosis).

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FEATURES OF PROTEIN ENCODED BY GENE NO: 69

Gene shares homology with a human collagen protein. Preferred polypeptide fragments comprise the following amino acid sequence:

5 MPRKTSKCRQLLCSGASRNADTAARQSTCSSHRPPGKIPSLGPRRXPGCXSVP SSRGEQSTGSPAAPRCGRRDAHRGLPGGAAMTPGDTWASFNPRAGHSKSQGE GQESSGASRQDRHPVSHWVERQREAWGAPRSSSAGGVKVAATTEREPEFKIK TGKA (SEQ ID NO:527); CSGASRNADTAARQSTCSSHRPPGKIPSLGPRRXPG CXSVPSSRGEQSTGSPAAPRCGRRDAHRGLPGGAAMTPGDTWASFNPRAGHS (SEQ ID NO:528); QGEGQESSGASRQDRHPVSHWVERQREAWGAPRSSSAGG VKVAATTEREPEFKIKTGKA (SEQ ID NO:529) (See Accession No. 124886). Also preferred are polynucleotide fragments encoding these polypeptide fragments

This gene is expressed primarily in fetal heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 302 as residues: Pro-32 to Ser-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cadiovascular disorders (e.g. heart disease, restenosis, atherosclerosis, stroke, angina, thrombosis).

FEATURES OF PROTEIN ENCODED BY GENE NO: 70

The translation product of this gene shares sequence homology with a chicken single-strand DNA-binding protein. Preferred polypeptide fragments comprise the following amino acid sequence:

MSPRYPGGPRPPLRIPNQALGGVPGSQPLLPSGMDPTRQQGHPNMGGPMQRM TPPRGMVPLGPQNYGGAMRPPLNALGGPGMPGMNMGPGGGRPWPNPTNAN WO 98/54963

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SIPYSSASPGNYVGPPGGGGPPGTPIMPSPADSTNSGDNMYTLMNAVPPGPNR
PNFPMGPGSDGPMGGLGGMESHHMNGSLGSGDMDSISKNSPNNMSLSNQP
GTPRDDGEMGGNFLNPFQSESYSPSMTMSV (SEQ ID NO:530); MSPRYPGG
PRPPLRIPNQALGGVPGSQPLLPSGMDPTRQQGHPNMGGPMQRMTPPRGMVP
5 LGPQNYGGAMRPPLNALGGPGMPGMNMGPGGGRPWPNPTNANSIPYSSASP
GNY (SEQ ID. NO:531); LNALGGPGMPGMNMGPGGGRPWPNPTNANSIPYSS
ASPGNYVGPPGGGGPPGTPIMPSPADSTNSGDNMYTLMNAVPPGPN (SEQ ID
NO:532); GPMGGLGGMESHHMNGSLGSGDMDSISKNSPNNMSLSNQPGTPR
DDGEMGGNFLNPFQSESYSPSMTMSV (SEQ ID NO:533); TCEHSSEAKAFHDY
10 (SEQ ID NO:534). Also preferred are polynucleotide fragments encoding these
polypeptide fragments. (See Accession No. 1562534)

This gene is expressed primarily in placenta and to a lesser extent in the fetal heart and a variety of other tissues and cell types.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities, fetal deficiencies, and particularly of the cardiovascular system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of developmental abnormalities or fetal deficiencies, ovarian and other endometrial cancers, reproductive dysfunction, cardiovascular disorders, and pre-natal disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 71

This gene is expressed primarily in fetal liver and to a lesser extent in the breast and testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, liver disorders (including hepatoblastomas) and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic and reproductive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). The expression in testes and breast indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of endocrine and reproductive disorders (e.g. sperm maturation, milk production, testicular and breast cancers).

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 72

This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. W93595).

This gene is expressed primarily in smooth muscle and to a lesser extent in brain.

25 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiovascular and neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes 30 for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular and central nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene 35 expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of restenosis, atherosclerosis, stroke, angina, thrombosis, wound healing and other conditions of heart disease. In addition, the expression in brain would suggest that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of developmental, degenerative and behavioral conditions of the brain and nervous system (e.g. schizophrenia, depression, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia, paranoia, addictive behavior and sleep disorders).

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FEATURES OF PROTEIN ENCODED BY GENE NO: 73

Gene shares homology with human stromalin-2. Preferred polypeptide fragments comprise the following amino acid sequence:

QAFVLLSDLLLIFSPQMIVGGRDFLRPLVFFPEATLQSELASFLMDHVFIQPGDL
GSGA (SEQ ID NO:535); ACSYLLCNPEFTFFSRADFARSQLVDLLTDRFQQE
LEELLQVG (SEQ ID NO:536),QKQLSSLRDRMVAFCELCQSCLSDVDTEIQEQV
ST (SEQ ID NO:537); QVILPALTLVYFSILWTLTHISKSDAS (SEQ ID NO:538);
STHDLTRWELYEPCCQLLQKAVDTGXVPHQV (SEQ ID NO:539). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No.R65208) This gene maps to chromosome 7, and therefore, may be used as a marker in linkage analysis for chromosome 7 (See Accession No.D52585).

This gene is expressed primarily in the brain (infant brain, adult brain, pituitary, cerebellum, hippocampus, schizophrenic hypothalmus, amygdala).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental and neurodegenerative diseases of the brain and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those

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comprising a sequence shown in SEQ ID NO: 306 as residues: Thr-25 to Lys-36, Lys-55 to Ser-63.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and treatment of developmental, degenerative and behavioral conditions of the brain and nervous system (e.g. schizophrenia, depression, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia, paranoia, addictive behavior and sleep disorders).

FEATURES OF PROTEIN ENCODED BY GENE NO: 74

10 This gene is expressed primarily in the hypothalamus of a human suffering from schizophrenia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the CNS particularly schizophrenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS, such as schizophrenia expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 307 as residues: Gly-38 to Ala-44.

The tissue distribution indicates that the protein products of this gene are useful for the study, diagnosis and treatment of schizophrenia and other disorders involving the CNS.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 75

Preferred polypeptides of the invention comprise the following amino acid sequence encoded by this gene:

LAVSTSFICCADISTALPLGSSRPAPAPRHREHEHGHQARPPRLLXTSLMPLSTP AAAQLLWTQLTPMGGRPGGRHSPPTLHTGPRALPPGPPHPSLHVAALSLLR (SEQ ID NO:540). Polynucleotides encoding such polypeptides are also provided.

This gene is expressed primarily in endometrial tumor and to a lesser extent in amniotic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive and immune disorders particularly cancers of those systems. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 308 as residues: Ser-3 to Arg-9.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune and reproductive disorders particularly cancers of those systems.

FEATURES OF PROTEIN ENCODED BY GENE NO: 76

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This gene is expressed primarily in kidney cortex and to a lesser extent in early stage human brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, renal disorders such as renal cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the kidney expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 309 as residues: Gly-38 to Gly-45, Gly-47 to Gly-52, Pro-92 to Lys-110.

The tissue distribution indicates that the protein products of this gene are useful for study, treatment and diagnosis of renal diseases such as cancer of the kidney.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 77

This gene is expressed primarily in kidney medulla.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, metabolic and renal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and renal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study, treatment and diagnosis of metabolic and renal diseases and disorders.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 78

This gene is expressed in chronic synovitis and microvascular endothelium.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, arthritis and atherosclerosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular and skeletal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study, diagnosis and treatment of arthritic and other inflammatory diseases as well as cardiovascular diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 79

This gene is expressed in resting T-cells and activated monocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for the study and treatment of immune diseases such as inflammatory conditions.

FEATURES OF PROTEIN ENCODED BY GENE NO: 80

This gene is expressed in a variety of immune system tissues, e.g., neutrophils, T-cells, and TNF induced epithelial and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and vascular systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 313 as residues: Met-1 to Trp-6.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of infectious diseases, immune and vascular disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 81

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and other immune conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 82

This gene is expressed in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory and other immune conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 315 as residues: Ala-83 to Thr-91.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 83

This gene is expressed in human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and inflammatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the inflammatory and immune systems.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 84

This gene is expressed in human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the inflammatory and immune systems. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the inflammatory and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the immune and inflammatory systems.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 85

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune system diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and inflammatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of diseases of the inflammatory and immune systems.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 86

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune system disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the inflammatory and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 319 as residues: Met-1 to Gly-6, Gly-32 to Pro-43, Leu-55 to Gln-60.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the immune and inflammatory system.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 87

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In specific embodiments, polypeptides of the invention comprise the sequence: **EQVLALLWPRFELILEMNVQSVRSTDPQRLGGLDTRPHYITRRYAEFSSALVSIN** OTIPNERTMQLLGQLQVEVENFVLRVAAEFSSRKEQLVFLINNYDMMLGVLME RAADDSKEVESFQQLLNARTQEFIEELLSPPFGGLVAFVKEAEALIERGQAERLR GEEARVTQLIRGFGSSWKSSVESLSQDVMRSFTNFRNGTSIIQG (SEQ ID NO:541),ALLKYRFFYQFLLGNERATAKEIRDEYVETLSKIYLSYYRSYLGRLMK VOYEEVAEKDDLMGVEDTAKKGFXSKPSRSRNTIFTLGTRGSVISPTELEAPILV PHTAQR (SEQ ID NO: 542); EQRYPFEALFRSQHYXLLDNSCREYLFICEFFVVS **GPXAHDLFHAVMGRTLSMTLKHLDSYLADCYDAIAVFLCIHIVLRFRNIAAKRD** VPALDRYW (SEQ ID NO:543),GGLDTRPHYITRRYAEFSSALVSINQ (SEQ ID NO:544); SRKEQLVFLINNYDMMLGVL (SEQ ID NO: 545) and/or ALLKYRFFY QFLLGNERATAKEIRDEYVETLSKIYLSYYRSYLGRLMKVQYEEVAEKDDLMG VEDTAKKGFXSKPSLRSRNTIFTLGTRGSVISPTELEAPILVPHTAQRXEQRYPF EALFRSQHYXLLDNSCREYLFICEFFVVSGPXAHDLFHAVMGRTLSMTLKHLD SYLADCYDAIAVFLCIHIVLRFRNIAAKRDVPALDRYWEQVLALLWPRFELILEM NVQSVRSTDPQRLGGLDTRPHYTTRRYAEFSSALVSINQTIPNERTMQLLGQLQV EVENFVLRVAAEFSSRKEQLVFLINNYDMMLGVLMERAADDSKEVESFQQLLN ARTQEFIEELLSPPFGGLVAFVKEAEALIERGQAERLRGEEARVTQLIRGFGSSW KSSVESLSQDVMRSFTNFRNGTS (SEQ ID NO:546). Polynucleotides encoding these polypeptides are also encompassed by the invention. The translation product of this gene shares sequence homology with suppressor of actin mutation which is thought to be important in mutation suppression.

This gene is expressed primarily in fetal liver and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver and mutations. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver or cancer, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level

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in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 320 as residues: Val-53 to Arg-60, Thr-88 to Thr-94, Ala-142 to Ser-150, Gly-188 to Glu-196, Gly-208 to Ser-214, Thr-227 to Gly-232, Lys-279 to Phe-285.

The tissue distribution and homology to suppressor of actin mutation suggest that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and of liver disorder or cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 88

This gene maps to chromosome 9, and therefore can be used in linkage analysis as a marker for chromosome 9. In specific embodiments, polypeptides of the invention comprise the sequence:

YEGKEFDYVFSIDVNEGGPSYKLPYNTSDDPWLTAYNFLQKNDLNPMFLDQVA KFIIDNTKGQMLGLGNPSFSDPFTGGGRYVPGSSGSSNTLPTADPFTGAGRYV PGSASMGTTMAGVDPFTGNSAYRSAASKTMNIYFPKKEAVTFDQANPTQILGK LKELNGTAPEEKKLTEDDLILLEKILSLICNSSSEKPTVQQLQILWKAINCPEDIV FPALDILRLSIKHPSVNENFCNEKEGAQFSSHLINLLNPKGKPANQLLALRTFC NCFVGQAGQKLMMSQRESLMSHAIELKSGSNKNI (SEQ ID NO: 547); HIALATLALNYSVCFHKD (SEQ ID NO: 548); HNIEGKAQCLSLISTILEVVO

- 20 DLEATFRLLVALGTLISDDSNAVQLAKS (SEQ ID NO:549); LGVDSQIKKYSS VSEPAKVSECCRFILNLL (SEQ ID NO:550); and/or YEGKEFDYVFSIDVNEGGPS YKLPYNTSDDPWLTAYNFLQKNDLNPMFLDQVAKFIIDNTKGQMLGLGNPSFS DPFTGGGRYVPGSSGSSNTLPTADPFTGAGRYVPGSASMGTTMAGVDPFTGN SAYRSAASKTMNIYFPKKEAVTFDQANPTQILGKLKELNGTAPEEKKLTEDDLI
- 25 LLEKILSLICNSSSEKPTVQQLQILWKAINCPEDIVFPALDILRLSIKHPSVNENFC
 NEKEGAQFSSHLINLLNPKGKPANQLLALRTFCNCFVGQAGQKLMMSQRESL
 MSHAIELKSGSNKNIHIALATLALNYSVCFHKDHNIEGKAQCLSLISTILEVVQD
 LEATFRLLVALGTLISDDSNAVQLAKSLGVDSQIKKYSSVSEPAKVSECCRFILN
 LL (SEQ ID NO:551). Polynucleotides encoding these polypeptides are also
 30 encompassed by the invention. These polypeptides share significant homeless which
 - encompassed by the invention. These polypeptides share significant homology with phospholipase A2 activating protein which is thought to be important in signal transduction (see, e.g., Wang et al., Gene 161(2):237-241 (1995)).

This gene is expressed primarily in endothelial cells, to a less extent in placenta, endometrial stromal cells, osteosarcoma, testis tumor, muscle, and infant brain that are likely to be rich in blood vessles.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in vascular system, aberrent angiogenesis, tumor angiogenesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system or tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene in endothelial cells and several potential highly vascularized tissues and its homology to phospholipase A2 activating protein suggest that this gene may be involved in transducing signals for endothelial cells in angiogenesis or vasculogenesis.

FEATURES OF PROTEIN ENCODED BY GENE NO: 89

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In specific embodiments, polypeptides of the invention comprise the sequence: YPNQDGDILRDQVLHEHIQRLSKVVTANHRALQIPEVYLREAPWPSAQSEIRTIS AYKTPRDKVQCILRMCSTIMNLLSLANEDSVPGADDFVPVLVFVLIKANPPCLL STVQYISSFYASCLSGEESYWWMQFTAAVE (SEQ ID NO:552); YPNQDGDILR DQVLHEHIQRLSKVVTANHRALQIPEVYLREAPWPSAQSEIRTISAYKTPRDKVQ CILRMCSTIMNLLSLANEDSVPGADDFVPVLVFVLIKANPPCLLSTVQYISSFYA SCLSGEESYWWMQFTAAVEFIKTI (SEQ ID NO:553); YPNQDGDILRDQVL (SEQ ID NO:554); EAPWPSAQSEI (SEQ ID NO:555); PVLVFVLIKANP (SEQ ID NO:560); SGEESYWWMQFTAAVEFIKTI (SEQ ID NO:556); ADDFVPVLVF VLIKANPP (SEQ ID NO:557); YKTPRDKVQCIL (SEQ ID NO:558); and/or GADDFVPVLVFVLIK (SEQ ID NO:559). The translation product of this gene shares sequence homology with human ras inhibitor and yeast VPS9p which is thought to be important in golgi vacuole transport.

This gene is expressed primarily in T cells and melanocytes and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, dysfunction and disorders involving T cells and melanocytes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing

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immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ras inhibitor indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating signal transduction; diagnosis and treatment of disorders involving T cells and melanocytes.

FEATURES OF PROTEIN ENCODED BY GENE NO: 90

This gene maps to chromosome 9 and therefore polypeptides of the invention can be used in linkage analysis as a marker for chromosome 9. The translation product of this gene shares sequence homology with neuronal olfactomedin-related ER localized protein which is thought to be important in influence the maintenance, growth, or differentiation of chemosensory cilia on the apical dendrites of olfactory neurons. In specific embodiments, polypeptides of the invention comprise the sequence: SARASTQPPAGQHPGPC (SEQ ID NO:561); MPGRWRWQRDMHPARKLLSLL FLILMGTELTQD (SEQ ID NO:562); SAAPDSLLRSSKGSTRGSL (SEQ ID NO:563); AAIVIWRGKSESRIAKTPGI (SEQ ID NO:564); FRGGGTLVLPPTHT PEWLIL (SEQ ID NO:567); PLGITLPLGAPETGGGD (SEQ ID NO:565); and/or CAAETWKGSQRAGQLCALLA (SEQ ID NO:566).

This gene is expressed in pineal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological and endocrinological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological or endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 323 as residues: Leu-20 to Ala-26, Arg-32 to Arg-39, Thr-104 to Gly-112.

The tissue distribution and homology to olfactomedin-related protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for maintenance, growth, or differentiation of neuron cells in pineal gland, therefore, may be useful for diagnosis and treatment of neurological disorders in pineal gland.

FEATURES OF PROTEIN ENCODED BY GENE NO: 91

This gene is expressed primarily in prostate and apoptotic T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate disease and T cell dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate cancer, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detect abnormal activity in prostate and T cells or probably treatment of this abnormality.

FEATURES OF PROTEIN ENCODED BY GENE NO: 92

This gene is expressed primarily in prostate and to a lesser extent in smooth muscle cells, fibroblasts, and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in prostate or vascular system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prosate or vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain

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tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating function of prostate or highly vascularized tissues, e.g. placenta.

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 93

This gene is expressed primarily in embryos and fetal tissues stage human and to a lesser extent in a wide variety of other proliferative tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in embryonic development and cell proliferation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the embryonic tissues and proliferative cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of abnormalities in developing and proliferative cells and organs.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 94

The translation product of this gene shares sequence homology with transformation related protein which is thought to be important in transformation.

This gene is expressed primarily in female reproductive tissues, i.e., breast cancer cells, placenta, and ovary and to a lesser extent in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, cancer or dysfunction of reproductive tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproduction system,

5 expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 327 as residues: Ser-50 to Pro-61.

The tissue distribution and homology to transformation related protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of conditions caused by transformation, i.e. tumorigenesis in reproductive organs, e.g. breast, placenta, and ovary.

FEATURES OF PROTEIN ENCODED BY GENE NO: 95

This gene is expressed primarily in testes, rhabdomyosarcoma, infant brain and to a lesser extent in some tumors and highly vascularized tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumorigenesis, abnormal angiogenesis, and/or neurological disorders., Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumor tissues or vascular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 328 as residues: Arg-46 to Trp-54, Pro-60 to Ile-69, Asn-116 to Ala-122, Arg-147 to Lys-153, Ser-158 to Glu-170, Ile-399 to Ser-405, Pro-486 to Met-499, Pro-502 to Asp-508.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for a range of disease states including treatment of

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tumor or vascular disorders and the treatment of neurological disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 96

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This gene maps to chromosome 7 and therefore polynucleotides of the present invention can be used in linkage analysis as a marker for chromosome 7. The translation product of this gene is homologous to the Clostridium perfringens enterotoxin (CPE) receptor gene product and shares sequence homology with a human ORF specific to prostate and a glycoprotein specific to oligodendrocytes both of which are tissue specific proteins.(See e.g., Katahira et al., J Cell Biol. 136(6):1239-1247 (1997). PMID: 9087440; UI: 97242441.

This gene is expressed primarily in pancreas tumor and ulcerative colitis and to a lesser extent in several tumors and normal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pancreatic disorder, ulcerative colitis, tumors and food poisoning. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system or tumorigenic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 329 as residues: Gly-147 to Met-152, Cys-177 to Lys-188.

The tissue distribution and homology to prostate and oligodendrocyte-specific protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for marker of diagnosis or treatment of disorder in pancreas, ulcerative colitis, and tumors. Furthermore, identity to the human receptor for Clostridium perfringenes entertoxin indicates that the soluble portion of this receptor could be used in the treatment of food poisoning associated with Clostridia perfringens by blocking the activity of perfringens enterotoxin.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 97

The translation product of this gene shares sequence homology with ATPase which is thought to be important in metabolism.

This gene is expressed primarily in testes and several hematopoietic cells and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, leukemia and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 330 as residues: Leu-37 to Ala-42.

The tissue distribution and homology to ATPase indicates that polynucleotides and polypeptides corresponding to this gene are useful for marker of diagnosis and treatment of leukemia and other hematopoietic disorders.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 98

In specific embodiments, polypeptides of the invention comprise the sequence: MRSARPSLGCLPSWAFSQALNI (SEQ ID NO:568); LLGLKGLAPAEISAVCE KGNFN (SEQ ID NO:569); VAHGLAWSYYIGYLRLILPELQARIR (SEQ ID NO:570); TYNQHYNNLLRGAVSQRC (SEQ ID NO:571); ILLPLDCGVPDNLSM ADPNIRFLDKLPQQTGDRAGIKDRVYSN (SEQ ID NO:572); SIYELLENGQRAGT CVLEYATPLQTLFAMSQYSQAGFSGEDRLEQ (SEQ ID NO:573); AKLFCRTLE DILADAPESQNNCRLIAYQEPADDSSFSLSQEVLRHLRQEEKEEVTVGSLKTSAV PSTSTMSQEPELLISGMEKPLPLRTDFS (SEQ ID NO:574); and/or LLGLKGLA PAEISAVCEKGNFNVAHGLAWSYYIGYLRLILPEL (SEQ ID NO:575).

35 Polynucleotides encoding these polypeptides are also encompassed by the invention.
This gene is expressed primarily in prostate BPH and to a lesser extent in bone marrow.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, benign prostatic hypertrophy or prostate cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male urinary system. expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 331 as residues: Ile-60 to Asn-69, Leu-106 to Asp-112, Glu-130 to Gly-136, Phe-160 to Glu-167, Pro-184 to Cys-190, Glu-197 to Ser-202, Arg-215 to Glu-221, Thr-237 to Pro-242.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of benign prostatic hypertrophy or prostate cancer.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 99

This gene is expressed primarily in salivary gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders or injuries of the salivary gland. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of glandular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of disorders of, or injuries to the salivary gland or other glandular tissue.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 100

This gene maps to chromosome 15, accordingly, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 15. The translation product of this gene shares sequence homology with a *C.elegans* gene of unknown function. In specific embodiments, polypeptides of the invention comprise the sequence: DPRVRLNSLTCKHIFISLTQ (SEQ ID NO:583); TMKLLKLRRNIV KLSLYRHFTN (SEQ ID NO:576); TLILAVAASIVFIIWTTMKFRI (SEQ ID NO:577); VTCQSDWRELWVDDAIWRLLFSMILFVI (SEQ ID NO:578); MVLWR PSANNQRFAFSPLSEEEEEDEQ (SEQ ID NO:580); KEPMLKESFEGMKMRS TKQEPNGNSKVNKAQEDDL (SEQ ID NO:584); and/or KWVEENVPSSVTDVALP ALLDSDEERMITHFERSKME (SEQ ID NO:582). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in thyroid and to a lesser extent in osteoclastoma, kidney medulla, and lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, thyroid dysfunction or cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 333 as residues: Lys-107 to Leu-124, Glu-150 to Thr-159, Pro-173 to Asp-179, Ser-192 to Ser-201.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of thyroid dysfunction or cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 101

This gene maps to chromosome 16, therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 16. In specific embodiments, polypeptides of the invention comprise the sequence:

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IRHELTVLRDTRPACA (SEQ ID NO:585); and/or MDFXMALIYD (SEQ ID NO:586). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in kidney cortex and to a lesser extent in adult brain, corpus colosum, hippocampus, and frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment or diagnosis of neurological disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 102

In specific embodiments, polypeptides of the invention comprise the sequence: MQEMMRNQDRALSNLESIPGGYNA (SEQ ID NO:587); LRRMYTDIQEPMLSA 25 AQEQF GGNPF (SEQ ID NO:588); ASLVSNTSSGEGSQPSRTENRDPLPNPWAP QT (SEQ ID NO:589); SQSSSASSGTASTVGGTTGSTASGTSGQSTTAPNLVPGV GASMFNTPG MQSLLQQITENPQLMQNMLSAPY (SEQ ID NO:590): MRSMMQSLSQNPDLAAQMMLNNPLFAGNPQLQEQMRQQLPTFLQQ (SEQ ID NO:591); MQNPDTLSAMSNPRAMQALLQIQQGLQTLATEAPGLIPGFTPGLG 30 ALGSTGGSSGTNGSNATPSENTSPTAGT (SEQ ID NO:592); TEPGHQOFI QQMLQALAGVNPQLQNPEVRFQQQLEQLSAMGFLNREANLQALIATGGDINAA IERLLGSQPS (SEQ ID NO:593); RNPAMMQEMMRNQDRALSNLESIPGGY NALRRMYTDIQEPMLSAA (SEQ ID NO:594); GNPFASLVSNTSS (SEQ ID NO:595); ENRDPLPNPWA (SEQ ID NO:595); GKILKDQDTLSQHGIHD (SEQ ID NO:597); GLTVHLVIKTQNRP (SEQ ID NO:598); SELQSQMQRQLLSNPEMM 35 (SEQ ID NO:599); PEISHMLNNPDIMR (SEQ ID NO:600); and/or RQLIMANPQMQQLIQRNP (SEQ ID NO:601). Polynucleotides encoding these

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polypeptides are also encompassed by the invention.

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This gene is expressed primarily in breast.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, breast cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of tumor systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of some types of breast cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 103

The translation product of this gene shares sequence homology with secreted serine proteases and lysozyme C precursor, which is thought to be important in bacteriolytic function. In specific embodiments, polypeptides of the invention comprise the sequence: NLCHVDCQDLLNPNLLAGIHCAKRIVS (SEQ ID NO:602); LDGFEGYSLSDWLCLAFVESKFN (SEQ ID NO:603);

25 NENADGSFDYGLFQINSHYWCN (SEQ ID NO:604); and/or NLCHVDCQDLLNPNLLAGIHCAKRIVS (SEQ ID NO:605). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infection. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 336 as residues: Ile-62 to Phe-70, Asn-78 to Asn-84.

The tissue distribution and homology to lysozyme C precursor indicates that polynucleotides and polypeptides corresponding to this gene are useful for boosting the moncyte-macrophage system and enhance the activity of immunoagents.

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 104

This gene is expressed primarily in apoptotic T-cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of some immune disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 105

The translation product of this gene shares sequence homology with ARI

protein of Drosophila (accession 2058299; EMBL: locus DMARIADNE, accession X98309), which is thought to be important in axonal path-finding in the central nervous system. In specific embodiments, polypeptides of the invention comprise the sequence IREVNEVIQNPAT (SEQ ID NO:606); ITRILLSHFNWDKEKLMERYF DGNLEKLFA (SEQ ID NO:607); NTRSSAQDMPCQICYLNYPNSYF (SEQ ID NO:608); TGLECGHKFCMQCWSEYLTTKIMEEGMGQTISCPAHG (SEQ ID NO:614); CDILVDDNTVMRLITDSKVKLKYQHLITNSFVECNRLLKWCPAPD CHHVVKVQYPDAKPV (SEQ ID NO:609); CDILVDDNTVMRLITDSK

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VKLKYQHLITNSFVECNRLLKWCPAPDCHHVVKV (SEQ ID NO:610);
GCNHMVCRNQNCKAEFCWVCLGPWEPHGSAWYNCNRYNEDDAKAARDAQE
RSRAALQRYL (SEQ ID NO:611); FYCNRYMNHMQSLRFEHKLYAQVKQ
KMEEMQQHNMSWIEVQFLKKAVDVLCQCRATLMYT (SEQ ID NO: 612);
YVFAFYLKKNNQSIIFENNQADLENATEVLSGYLERDISQDSLQDIKQKVQDKY
RYCESR (SEQ ID NO:613) Polynucleotides encoding these polypeptides are also
encompassed by the invention.

This gene is expressed primarily in adult brain, and to a lesser extent in endometrial tumor, melanocytes, and infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases or injuries involving axonal path development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ARI protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of disease states or injuries involving axonal path development, including neurodegenerative diseases and nerve injury.

FEATURES OF PROTEIN ENCODED BY GENE NO: 106

The translation product of this gene shares sequence homology with cytochrome b561 [Sus scrofa] which is thought to be an integral membrane protein of neuroendocrine storage vesicles of neurotransmitters and peptide hormones.

This gene is expressed primarily in frontal cortex and to a lesser extent in rhabdomyosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to

these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 339 as residues: Ser-18 to Pro-24.

The tissue distribution and homology to cytochrome b561 [Sus scrofa] indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of neurological disorders. This gene may also be important in regulation of some types of cancers.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 107

In specific embodiments, polypeptides of the invention comprise the sequence: MWGYLFVDAAWNFLGCLICGW (SEQ ID NO:615); MHFISSGNVSAIRSSILLL RXSLSYLGNCLRVSAIFVYFLLFLLLS (SEQ ID NO:616); and/or MDQALRGSPSE GFSTDPSPPQVGRQIPSFPPWRRLVLPKASGCFLEREWWLCVFKLRTRPGAEA HAYNSSILGGRGKGIT (SEQ ID NO:617). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in pancreas tumor and to a lesser extent in cerebellum.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pancreatic tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred

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epitopes include those comprising a sequence shown in SEQ ID NO: 340 as residues: Pro-22 to Phe-33.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of pancreatic tumors.

FEATURES OF PROTEIN ENCODED BY GENE NO: 108

This gene maps to chromosome 17 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 17. In specific embodiments, polypeptides of the invention comprise the sequence:

MLPALASCCHFSPPEQAARLKKLQEQEKQQKVEFRKRMEKEVSDFIQDSGQIK KKFQPMNKIERSILHDVVEVAGLTSFSFGEDDDCRYVMIFKKEFAPSDEELDSY RRGEEWDPQKAEEKRNXKELAQRQ (SEQ ID NO:618); EEEAAQQGPVVV SPASDYKDKYSHLIGKGAAKDAAHMLQANKTYGCXPVANKRDTRSIEEAMNE IRAKKRLRQSGE (SEQ ID NO:619); PPRRPAQLPLTPGAGQGAGRDKAAAIRA HPGAPPLNHLLP (SEQ IDNO:620); AVPQAGGKQVFDLSPLELGYVRGMCVCV (SEQ ID NO:621) and/or MLPALASCCHFSPPEQAARLKKLQEQEKQQKVEFRK RMEKEVSDFIQDSGQIKKKFQPMNKIERSILHDVVEVAGLTSFSFGEDDDCRYV MIFKKEFAPSDEELDSYRRGEEWDPQKAEEKRNXKELAQRQEEEAAQQGPVVV SPASDYKDKYSHLIGKGAAKDAAHMLQANKTYGCXPVANKRDTRSIEEAMNE

20 IRAKKRLRQSGE (SEQ ID NO:622). Polynucleotides encoding these polypeptides are also encompassed by the invention. The translation product of this gene shares sequence homology with FSA-1 which may play a role as a structural protein component of the acrosome.

This gene is expressed primarily in fetal kidney and sperm.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders, especially involving acrosomal disfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

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individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 341 as residues: Glu-8 to Asn-35.

The tissue distribution and homology to FSA-1 indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of infertility due to acrosomal disfunction of sperm.

FEATURES OF PROTEIN ENCODED BY GENE NO: 109

This gene is expressed primarily in pituitary and to a lesser extent in epididymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 342 as residues: Met-1 to Trp-6.

Because the gene is found in both pituitary and epididymus, this indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of male reproductive disorders. This may involve a secreted peptide produced in the pituitary targeting the epididymus.

FEATURES OF PROTEIN ENCODED BY GENE NO: 110

In specific embodiments, polypeptides of the invention comprise the sequence: LLCPVLNSGXSWNFPHPSQPEYSFHGFHSTRLWI (SEQ ID NO:623); and/or PSTPWFLFLLGLTCPFSTSHPRWDSIPP (SEQ ID NO:624). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in resting T-cells. .

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, T-cell disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of certain immune disorders, especially those involving T-cells.

FEATURES OF PROTEIN ENCODED BY GENE NO: 111

This gene is expressed primarily in cerebellum and whole brain and to a lesser extent in infant brain and fetal kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 344 as residues: Asp-48 to Gly-55.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological disorders.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 112

The translation product of this gene shares sequence homology with yeast mitochondrial ribosomal protein homologous to ribosomal protein s15 of E.coli which

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is thought to be important in the early assembly of ribosomes (See Accession No. M38016). This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1.

This gene is expressed primarily in developmental tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development of cancers and tumors in addition to healing wounds. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and developmental expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ribosomalprotein s15 of E. coli indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases related to the assembly of ribosomes in the mitochondria which is important in the translation of RNA into protein. Therefore, this indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of multiple tumors as well as in healing wounds which are thought to be under similar regulation as developmental tissues. Protein, as well as, antibodies directed against the protein have utility as tumor markers, in addition to immunotherapy targets, for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 113

The translation product of this gene shares sequence homology with human poliovirus receptor precursors which are thought to be important in viral binding and uptake. Preferred polypeptide fragments comprise the following amino acid sequence: ELSISISNVALADEGEYTCSIFTMPVRTAKSLVTVLGIPQKPIITGYKSSLREKDT ATLNCQSSGSKPAARLTWRKGDQELHGEPTRIQEDPNGKTFTVSSSVTFQVTR EDDGASIVCSVNHESLKGADRSTSQRIEVLYTPTAMIRPDPPHPREGQKLLLHC EGRGNPVPQQYLWEKEGSVPPLKMTQESALIFPFLNKSDSGTYGCTATSNMGS YKAYYTLNVND (SEQ ID NO:625). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. gnllPIDId1002627).

This gene is expressed almost exclusively in human brain tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, susceptibility to viral disease and diseases of the CNS especially cancers of that system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 346 as residues: Leu-26 to Asp-37, Lys-53 to Ser-59.

The tissue distribution and homology to poliovirus receptor precursors indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and prevention of diseases that involve the binding and uptake of virus particles for infection. It might also be helpful in genetic therapy where the goal is to insert foreign DNA into infected cells. With the help of this protein, the binding and uptake of this foreign DNA might be aided. In addition, it is expected that over expression of this gene will indicate abnormalities involving the CNS, particularly cancers of that system.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 114

The translation product of this gene shares sequence homology with YO87_CAEEL hypothetical 28.5 KD protein ZK1236.7 in chromosome III of Caenorhabditis elegans in addition to alpha-1 collagen type III (See Accession No. gil537432). One embodiment for this gene is the polypeptide fragment(s) comprising the following amino acid sequence: VPELPDRVHQLHQAVQGCALGRPGFPGGPTH SGHHKSHPGPAGGDYNRCDRPGQVHLHNPRGTGRRGQLHPTAGPGVHRRA CPSQQLPHRLGPGVPCPSPSLTPVLPSWTQSWCG LPGYTSSS (SEQ ID NO:630). An additional embodiment is the polynucleotide fragment(s) encoding these polypeptide fragments

This gene is expressed primarily in brain cells and to a lesser extent in activated B and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegeneration and imunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neural and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 347 as residues: Glu-34 to Glu-39, Gly-51 to Ser-72, Ala-88 to Glu-93, Gln-100 to Val-105.

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The tissue distribution and homology to YO87_CAEEL hypothetical 28.5 KD protein ZK1236.7 in chromosome III of Caenorhabditis elegans as well as to a conserved alpha-1 collagen type III protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons' Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorders. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 115

The translation product of this gene shares sequence homology with alpha 3 type IX collagen which is thought to be important in hyaline cartilage formation via its ability to uptake inorganic sulfate by cells (See Accession No. gil975657). One embodiment of this gene is the polypeptide fragment comprising the following amino acid sequence: SLRRPRSAAXQTLTTFLSSVSSASSSALPGSREPCDPRAPPPPR SGSAASCCSCCCSCPRRRAPLRSPRGSKRRIRQREVVDLYNGMCLQGPAGVPG RDGSPGANGIPGTPGIPGRDGFKGEKGECLRESFEESWTPNYKQCSWSSLNY GIDLGKIAECTFTKMRSNSALRVLFSGSLRLKCRNACCQRWYFTFNGAECSGP LPIEAIIYLDQGSPEMNSTINIHRTSSVEGLCEGIGAGLVDVAIWVGTCSDYPKG DASTGWNSVSRIIIEELPK (SEQ ID NO:634). An additional embodiment are the

polynucleotide fragments encoding this polypeptide fragment.

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This gene is expressed primarily in smooth muscle and to a lesser extent in synovial tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, dwarfism, spinal deformation, and specific joint abnormalities as well as chondrodysplasias i.e., spondyloepiphyseal dysplasia congenita, familial osteoarthritis, Atelosteogenesis type II, metaphyseal chondrodysplasia type Schmid and autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to alpha 3 type IX collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of diseases associated with the mutation in this gene which leads to the many different types of chondrodysplasias. By the use of this product, the abnormal growth and development of bones of the limbs and spine could be routinely detected or treated in utero since the protein or muteins thereof could affect epithelial cells early in development and later the chondrocytes of the developing craniofacial structure.

FEATURES OF PROTEIN ENCODED BY GENE NO: 116

The translation product of this gene shares sequence homology with retrovirus-related reverse transcriptase which is thought to be important in viral replication. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: TKKENCRPASLMNIDTKILNKILMNQ (SEQ ID NO:640). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments (See Accession No. pirlA25313IGNHUL1).

This gene is expressed primarily in human meningima.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, retroviral diseases such as AIDS, and possibly certain cancers due to transactivation of latent cell division genes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to retrovirus-related reverse transcriptase indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of diseases and maladies associated with retroviral infection since a functional reverse transcriptase (RT) or RT-like molecule is an integral component of the retroviral life cycle.

FEATURES OF PROTEIN ENCODED BY GENE NO: 117

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The translation product of this gene shares sequence homology with an unknown gene from *C. elegans*, as well as weak homolog with mammalian metaxin, a gene contiguous to both thrombospondin 3 and glucocerebrosidase, is known to be required for embryonic development. Preferred polypeptide fragments comprise the following amino acid sequence: MCNLPIKVVCRANAEYMSPSGKVPXXHVGNQ VVSELGPIVQFVKAKGHSLSDGLEEVQKAEMKAYMELVNNMLLTAELYLQWC DEATVGXITHXRYGSPYPWPLXHILAYQKQWEVKRKXKAIGWGKKTLDQVLE DVDQCCQALSQRLGTQPYFFNKQPTELDALVFGHLYTILTTQLTNDELSEKVKN YSNLLAFCRRI EQHYFEDRGKGRLS (SEQ ID NO:641); MCNLPIKVVCRANAE YMSPSGKVPXXHVGNQVVSELGPIVQFVK (SEQ ID NO:642),. Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. gil1326108).

This gene is expressed primarily in fetal tissues and to a lesser extent in hematopoietic cells and tissues, including spleen, monocytes, and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer; lymphoproliferative disorders; inflammation; chondrosarcoma, and Gaucher disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification

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of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and embryonic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of cancer and other proliferative disorders. Expression in embryonic tissue and other cellular sources marked by proliferating cells indicates that this protein may play a role in the regulation or cellular division. Additionally, the expression in hematopoietic cells and tissues indicates that this protein may play a role in the proliferation, differentiation, and survival of hematopoietic cell lineages. Thus, this gene may be useful in the treatment of lymphoproliferative disorders, and in the maintenance and differentiation of various hematopoietic lineages from early hematopoietic stem and committed progenitor cells.

FEATURES OF PROTEIN ENCODED BY GENE NO: 118

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The translation product of this gene shares sequence homology with reverse transcriptase which is important in the synthesis of a cDNA chain from an RNA molecule, and is a method whereby the infecting RNA chains of retroviruses are transcribed into their DNA complements. One embodiment for this gene is the polypeptide fragment comprising the following amino acid sequence:

MXXXNSHITIFTLNVNGLNAPNERHRLANWIQSQDQVCCIQETHLTGRDTHRL KIKGWRKIYQANGKQKK (SEQ ID NO:647). An additional embodiment is the polynucleotide fragments comprising polynucleotides encoding these polypeptide fragments (See Accession No. gil2072964).

This gene is expressed primarily in skin and to a lesser extent in neutrophils. Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer, hematopoietic disorders; inflammation; disorders of immune surveillance. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the epidermis and/or hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and

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wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to reverse transcriptase indicates that polynucleotides and polypeptides corresponding to this gene are useful for cancer therapy. Expression in the skin also indicates that this gene is useful in wound healing and fibrosis. Expression by neutrophils also indicates that this gene product plays a role in inflammation and the control of immune surveillance (i.e. recognition of viral pathogens). Reverse transcriptase family members are also useful in the detection and treatment of AIDS.

FEATURES OF PROTEIN ENCODED BY GENE NO: 119

The translation product of this gene shares sequence homology with reverse transcriptase which is important in the synthesis of a cDNA copy of an RNA molecule, and is a method whereby a retrovirus reverse-transcribes its genome into an inheritable DNA copy.

This gene is expressed primarily in the frontal cortex of brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer and neurodegenerative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS and peripheral nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to reverse transcriptase suggest that this is useful in the treatment of cancer and AIDS. The expression in brain indicates that it plays a role in neurodegenerative disorders and in neural degeneration.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 120

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One embodiment of this gene has homology to a hypothetical protein in Schizosaccharomyces pombe (See Accession No. 2281980). Another embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: IYHLHSWIFFHFKRAFCMCFITMKVIHAHCSKLRKCXNAQISVFCTTLTASYPT (SEQ ID NO:651). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments. This gene maps to chromosome 18, and therefore, may be used as a marker in linkage analysis for chromosome 18.

This gene is expressed primarily in adult hypothalamus and to a lesser extent in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disorders; endocrine function; and vertigo. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, CNS and peripheral nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of neurodegenerative disorders; diagnosis of tumors of a brain or neuronal origin; treatments involving hormonal control of the entire body and of homeostasis, behavioral disorders, such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo.

FEATURES OF PROTEIN ENCODED BY GENE NO: 121

The translation product of this gene shares sequence homology with the human IRLB protein which is thought to be important in binding to a c-myc promoter element and thus regulating its transcription (See Accession No. gil33969). This gene maps to

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chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1.

This gene is expressed primarily in brain and breast and to a lesser extent in a variety of hematopoietic tissues and cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer of the brain and breast; lymphoproliferative disorders; neurodegenerative diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS, breast, and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cancer of the brain, breast, and hematopoietic system. In addition, it may be useful for the treatment of neurodegenerative disorders, as well as disorders of the hematopoietic system, including defects in immune competency and inflammation. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 122

The translation product of this gene shares sequence homology with an ATP synthase, a key component of the proton channel that is thought to be important in the translocation of protons across the membrane.

This gene is expressed primarily in T cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, T cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or

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lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ATP synthase indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of defects in proton transport, homeostasis, and metabolism, as well as the diagnosis and treatment of lymphoma. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia

FEATURES OF PROTEIN ENCODED BY GENE NO: 123

This gene maps to chromosome 15, and therefore, may be used as a marker in linkage analysis for chromosome 15.

This gene is expressed primarily in a variety of fetal tissues, including fetal liver, lung, and spleen, and to a lesser extent in a variety of blood cells, including eosinophils and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer (abnormal cell proliferation); T cell lymphomas; and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetus and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of conditions involving cell proliferation. Expression of this gene in fetal tissues, as well as in a variety of blood cell lineages indicates that it may play a role in either cellular proliferation; apoptosis; or cell survival. Thus it may be useful in the management and

treatment of a variety of cancers and malignancies. In addition, its expression in blood cells suggest that it may play additional roles in hematopoietic disorders and conditions, and could be useful in treating diseases involving autoimmunity, immune modulation, immune surveillance, and inflammation..

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FEATURES OF PROTEIN ENCODED BY GENE NO: 124

This gene is expressed primarily in placenta and to a lesser extent in pineal gland and rhabdomyosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental, endocrine, and female reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the [insert system where a related disease state is likely, e.g., immune], expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 357 as residues: Leu-69 to Val-76.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders in development. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 125

This gene is expressed primarily in benign prostatic hyperplasia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of benign prostatic hyperplasia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive

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system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of benign prostatic hyperplasia. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 126

This gene is expressed primarily in apoptotic T-cells and to a lesser extent in suppressor T cells and ulcerative colitis.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases involving premature apoptosis, and immunological and gastrointestinal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of disorders involving inappropriate levels of apoptosis, especially in immune cell lineages. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases (such as AIDS), and leukemia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 127

This gene is expressed primarily in Raji cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and T cell autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 360 as residues: Asp-23 to Gly-29.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of inflammation and T cell autoimmune disorders. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases (such as AIDS), and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 128

The translation product of this gene shares sequence homology with an *C. elegans* coding region C47D12.2 of unknown function (See Accession No. gnllPIDle348986). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: EDDGFNRSIHEVILKNITWY SERVLTEISLGSLLILVVIRTIQYNMTRTRDKYLHTNCLAALANMSAQFRSLHQY AAQRIISLFSLLSKKHNKVLEQATQSLRGSLSSNDVPLPDYAQDLNVIEEVIRMM LEIINSCLTNSLHHNPNLVALLYKRDLFEQFRTHPSFQDIMQNIDLVISFFSSRLL QAGS (SEQ ID NO:657); EDDGFNRSIHEVILKNITWYSERVLTEISLGSLLILVV (SEQ ID NO:658); RTIQYNMTRTRDKYLHTNCLAALANMSAQFRSLHQYAAQ RIISLFSLLSKKHN (SEQ ID NO:659); KKHNKVLEQATQSLRGSLSSNDVPLPDY AQD (SEQ ID NO:661); SCLTNSLHHNPNLVYALLYKRDLFEQFRTHPSFQD IMQNIDLVISFFSSRLLQAGS (SEQ ID NO:660). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments. This gene maps to

chromosome 18, and therefore, may be used as a marker in linkage analysis for chromosome 18.

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This gene is expressed primarily in smooth muscle and to a lesser extent in fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, atherosclerosis and other cardiovascular and hepatic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of circulatory system disorders such as atherosclerosis, hypertension, and thrombosis. In addition, the tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

FEATURES OF PROTEIN ENCODED BY GENE NO: 129

The translation product of this gene shares sequence homology with a ribosomal protein which is thought to be important in cellular metabolism, in addition to the C.elegans protein F40F11.1 which does not have a known function at the current time (See Accession No. gnllPIDle244552). Preferred polypeptide fragments comprise the following amino acid sequence:

35 MADIQTERAYOKOPTIFONKKRVLLGETGKEKLPRVTNKNIGLGFKDT PRRLLRGTYIDKKCPFTGNVSIRGRILSGVVTQDEDAEDHCHPPRLSALHPQVQ PLREAPQEHVCTPVPL LQGRPDR (SEQ ID NO:662); MKMQRTIVIRRDYLH

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YIRKYNRFEKRHKNMSVHLSPCFRDVQIGDIVTVGECRPLSKTVRFNVLKVTK AAGTKKQFQKF (SEQ ID NO:663); MADIQTERAYQKQPTIFQNKKRVLLGET GK (SEQ ID NO:664); HCHPPRLSALHPQVQPLREAPQEHVCTPVPL LQGRPDR (SEQ ID NO:666); NIGLGFKDTPRRLLRGTYIDKKCPFTGNVSIRGRILSGVVTQ (SEQ ID NO:669); MKMQRTIVIRRDYLHYIRKYNRFEKRHKNMSVHLSP (SEQ ID NO:667); CFRDVQIGDIVTVGECRPLSKTVRFNVLKVTKAAGTKKQFQKF (SEQ ID NO:668). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in Wilm's tumor and to a lesser extent in thymus and stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases affecting RNA translation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Wilm's tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 362 as residues: Thr-11 to Asp-20.

The tissue distribution and homology to a ribosomal protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases affecting RNA translation.

FEATURES OF PROTEIN ENCODED BY GENE NO: 130

The translation product of this gene shares sequence homology with a yeast DNA helicase which is thought to be important in global transcriptional regulation (See Accession No. gnllPIDle243594). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: IFYDSDWNPTVDQQA MDRAHRLGQTKQVTVYRLICKGTIEERILQRAKEKSEIQRMVISG (SEQ ID NO:670); TRMIDLLEEYMVYRKHTYXRLDGSSKISERRDMVADFQNRNDI FVFLLSTRAGGLGINLTAXDTVHF (SEQ ID NO:671); TRMIDLLEEYMVYRK HTYXRLDGSSKISERRDM (SEQ ID NO:674); RRDMVADFQNRNDIFVFLL

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STRAGGLGINLTAXDTVHF (SEQ ID NO:675), IFYDSDWNPTVDQQAMD RAHRLGQTKQVTVYRLICKG (SEQ ID NO:676); RLICKGTIEERILQRAK EKSEIQRMVISG (SEQ ID NO:678). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in amygdala.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases and disorders of the brain. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a DNA helicase indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases affecting RNA transcription, particularly developmental disorders and healing wounds since the later are though to approximate developmental transcriptional regulation.

FEATURES OF PROTEIN ENCODED BY GENE NO: 131

This gene is expressed primarily in prostate and to a lesser extent in amygdala and pancreatic tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate enlargement and gastrointestinal disorders, particularly of the pancreas and gall bladder. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to

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the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of prostate diseases, including benign prostatic hyperplasia and prostate cancer. In addition, the tissue distribution in tumors of the pancreas indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tissues where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 132

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This gene is expressed primarily in adult lung and to a lesser extent in hypothalamus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pulmonary diseases and neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the pulmonary and respiratory systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of pulmonary and respiratory disorders such as emphysema, pneumonia, and pulmonary edema and emboli. In addition, the tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental

disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 133

This gene is expressed primarily in human liver.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cirrhosis of the liver and other hepatic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of liver disorders such as cirrhosis, jaundice, and Hepatitus. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tissues.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 134

This gene is expressed primarily in fetal kidney and to a lesser extent in fetal liver and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development and regeneration of liver and kidney and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive and excretory systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 367 as residues: Pro-70 to Arg-77, Tyr-102 to Thr-107.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the kidney and liver, such as cirrhosis, kidney failure, kidney stones, and liver failure, hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells. In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 135

This gene is expressed primarily in brain, bone marrow, and to a lesser extent in placenta, T cell, testis and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative and immunological diseases and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 368 as residues: Met-1 to His-6.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also

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play a role in the treatment and/or detection of developmental disorders associated with the developing embryo, or sexually-linked disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 136

Translatation product of this gene is homologous to the human WD repeat protein HAN11. Preferred polypeptide fragments comprise the following amino acid sequence:

MSLHGKRKEIYKYEAPWTVYAMNWSVRPDKRFRLALGSFVEEYNNKVQLVG LDEESSEFICRNTFDHPYPTTKLMWIPDTKGVYPDLLATSGDYLRVWRVGETET RLECLLNNNKNSDFCAPLTSFDWNEVDPYLLGTSSIDTTCTIWGLETGQVLGRV NLVSGHVKTQLIAHDKEVYDIAFSRAGGGRDMFASVGADGSVRMFDLRHLEH STIIYEDPQHHPLLRLCWNKQDPNYLATMAMDGMEVVILDVRVPAHLXPGTTIE HVSMALLGPHIHPATSALQRMTTRLSSGTSSKCPEPLRTLSWPTQLXGEINNVQ WASTQPELSPSATTTAWRYSECSVGGAVPTRQGLLYFLPLPHPQS (SEQ ID

15 NO:679); MSLHGKRKEIYKYEAPWTVYAMNWSVRPDKRFRLALGSFV
EEYNNKVQLVGLDEESSEFICRNTFDHPYPTTKLMWIPDTKGVYPDLLATSGDY
LRVWRVGETETRLECLLNNNKNSDFCAPLTSFDWNEVDPYLL (SEQ ID
NO:680); SFDWNEVDPYLLGTSSIDTTCTIWGLETGQVLGRVNLVSGHVK
TQLIAHDKEVYDIAFSRAGGGRDMFASVGADGSVRMFDLRHLEHSTIIYEDPQH

20 HPLLRLCWNKQDPNYLATMAMDGMEVVILDVRVPAHLXPGTTI (SEQ ID NO:681); VGADGSVRMFDLRHLEHSTIIYEDPQHHPLLRLCWNKQDPNYLA TMAMDGMEVVILDVRVPAHLXPGTTIEHVSMALLGPHIHPATSALQRMTTRLS SGTSSKCPEPLRTLSWPTQLXGEINNVQWASTQPELSPSATTTAWRYSECSVG GAVPTRQGLLYFLPLPHPQS (SEQ ID NO:682). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in placenta, embryo, T cell and fetal lung and to a lesser extent in endothelial, tonsil and bone marrow.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological and developmental diseases in addition to cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or

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cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 369 as residues: Gly-19 to Gln-28, Pro-36 to Phe-42.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 137

This gene is expressed primarily in TNF and INF induced epithelial cells, T cells and kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory conditions particularly inflammatory reactions in the kidney. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 370 as residues: Thr-67 to Gly-72, Gln-132 to Ala-145, Arg-150 to Pro-157.

The tissue distribution indicates that the protein products of this gene are useful for treating the damage caused by inflammation of the kidney.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 138

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This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. D63485).

This gene is expressed primarily in breast cancer and colon cancer and to a lesser extent in thymus and fetal spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers, especially of the breast and colon tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 139

This gene maps to chromosome 17, and therefore, can be used as a marker for linkage analysis from chromosome 17.

This gene is expressed primarily in CD34 positive cells, and to lesser extent in activated T-cells and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunologically related diseases and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and hematopoietic system, expression of this gene at significantly higher or lower levels

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may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in CD34, T-cell and neutrophils indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of hematopoietic disorders and immunologically related diseases, such as anemia, leukemia, inflammation, infection, allergy, immunodeficiency disorders, arthritis, asthma, immune deficiency diseases such as AIDS.

FEATURES OF PROTEIN ENCODED BY GENE NO: 140

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This gene was recently cloned by another group, who called the gene KIAA0313 gene. (See Accession No. d1021609.) Preferred polypeptide fragments comprise the amino acid sequence:

- LYATATVISSPSTEXLSQDQGDRASLDAADSGRGSWTSCSSGSHDNIQTIQ
 HQRSWETLPFGHTHFDYSGDPAGLWASSSHMDQIMFSDHSTKYNRQNQSRES
 LEQAQSRASWASSTGYWGEDSEGDTGTIKRRGGKDVSIEAESSSLTSVTTEETK
 PVPMPAHIAVASSTTKGLIARKEGRYREPPPTPPGYIGIPITDFPEGHSHPARKP
 PDYNVALQRSRMVARSSDTAGPSSVQQPHGHPTSSRPVNKPQWHKXNESDPR
- 20 PDYNVALQRSRMVARSSDTAGPSSVQQPHGHPTSSRPVNKPQWHKXNESDP LAPYQSQGFSTEEDEDEQVSAV (SEQ ID NO:683); HMDQIMFSDHSTKYNRQ NQSRESLEQAQSRASWASSTGYWGE (SEQ ID NO:684); SVTTEETKPVPMP AHIAVASSTTKGLIARKEGRYREPPPTPPGYIGIPITD (SEQ ID NO:685); and VALQRSRMVARSSDTAGPSSVQQPHGHPTSSRPVNKPQW
- 25 HKXNESDPRLAPYQSQGF (SEQ ID NO:686). Also preferred are polynucleotide fragments encoding these polypeptide fragments. This gene maps to chromosome 4, and therefore, may be used as a marker in linkage analysis for chromosome 4 (See Accession No. AB002311).

This gene is expressed primarily in ovarian cancer, tumors of the Testis, brain, and colon.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, ovarian, testicle, brain and colon cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male and female reproductive systems,

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expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon, ovary, testis, and brain origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 141

This gene is expressed primarily in spleen and colon cancer.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, colon cancer and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal trace and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 142

Translation product is homologous to T cell translocation protein, a putative zinc finger factor (See Accession No. 340454), as well as to the G-protein coupled receptor TM5 consensus polypeptide (See Accession No. R50734). Preferred polypeptide fragments comprise the following amino acid sequence:

CLLFVFVSLGMRCLFWTIVYNVLYLKHKCNTVLLCYHLCSI (SEQ·ID NO:687); ACSKLIPAFEMVMRAKDNVYHLDCFACQLCNQRXCVGDKFFLKNNXXLCQT DYEEGLMKEGYAPXVR (SEQ ID NO:688). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in fetal brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders including brain cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Central Nervous System, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo.

FEATURES OF PROTEIN ENCODED BY GENE NO: 143

Translation product for this gene has significant homology to the Fas ligand, which is a cysteine-rich type II transmembrane protein/tumor necrosis factor receptor homolog. Mutations within this protein have been shown to result in generalized lymphoproliferative disease leading to the development of lymphadenopathy and autoimmune disease (See Medline Article No. 94185175). Preferred polypeptide

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fragments comprise the following amino acid sequence:
SALSEPGAPDRRRPCPESVPRRPDDEQWPPPTALCLDVAPLPPSS (SEQ ID NO:689). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. 473565).

This gene is expressed primarily in osteoblasts, lung, and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, osteoblast-related, pulmonary, neurological, and immunological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 376 as residues: Trp-33 to Thr-40, Lys-45 to Ile-63.

The tissue distribution in osteoblasts, lung, and brain combined with its homology to the Fas ligand indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the Fas ligand gene is known to be expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including asthma, immune deficiency diseases such as AIDS and leukemia, and various autoimmune disorders including lupus and arthritis.

FEATURES OF PROTEIN ENCODED BY GENE NO: 144

This gene shares sequence homology with a 21.5 KD transmembrane protein in the SEC15-SAP4 intergenic region of yeast. (See Accession No. 1723971.) Preferred polypeptide fragments comprise the amino acid sequence:

AHASESGERWWACCGVRFGLRSIEAIGRSCCHDGPGGLVANRGRRFKWAIEL SGPGGGSRGRSDRGSGQGDSLYPVGYLDKQVPDTSVQETDRILVEKRCWDIAL

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GPLKQIPMNLFIMYMAGNTISIFPTMMVCMMAWRPIQALMAISATFKMLESSSQ KFLQGLVYLIGNLMGLALAVYKCQSMGLLPTHASDWLAFIEPPERMEFSGG GLLL (SEQ ID NO:691); PVGYLDKQVPDTSVQETDRILVEKRCWDIALGPLKQ IPMNLFI (SEQ ID NO:693); and ATFKMLESSSQKFLQGLVYLIGNLMGLALAV YKCQSMGLLPTHASD (SEQ ID NO:692). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in osteoclastoma, hemangiopericytoma, liver, lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, osteoclastoma, hemangiopericytoma, liver and lung tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the above tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the lung and liver systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing osteoclastoma, hemangiopericytoma, liver and lung tumors.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 145

Translation product of this gene shares homology with the glucagon-69 gene which may indicate this gene plays a role in regulating metabolism. (See Accession No. A60318) One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

PTTKLDIMEKKKHIQIRFPSFYHKLVDSGRMRSKRETRREDSDTKHNL (SEQ ID NO:694). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in brain, kidney, colon, and testis.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, brain, kidney, colon, and testicular cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, neurological, circulatory, and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of brain, kidney, colon, and testis origins, indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 146

The translation product of this gene shares sequence homology with goliath protein which is thought to be important in the regulation of gene expression during development. Protein may serve as a transcription factor. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

- TEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMPPKNFSRGSLVFVSISFIV LMIISSAWLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTVKKGDKETD PDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCPMCKLNILKA LGIV (SEQ ID NO:695); TEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMP PKNFSRGSLVFVSISFIVLM IISSAWLIFYF (SEQ ID NO:697); SISFIVLMIISSA
- 35 WLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTVKKGDKE (SEQ ID NO:698); VKKGDKETDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDP

WLSEHCTCPMCKLNILKALGIV (SEQ ID NO:699). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments (See Accession No. 157535). Moreover, another embodiment is the polynucleotide fragments encoding these polypeptide fragments:

- 5 MTHPGTEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMPPKNFSRGS
 LVFVSISFIVLMIISSAWLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTV
 KKGDKETDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCP
 MCKLNILKALGIVPNLPCTDNVAFDMERLTRTQAVNRRSALGDLAGDNSLGLE
 PLRTSGISPLPQDGELTPRTGEINIAVTKEWFIIASFGLLSALTLCYMIIRATASLN
 10 ANEVEWF (SEQ ID NO:696);MTHPGTEHIIAVMITELRGKDILSYLEKNISVQM
 TIAVGTRMPPKNFSRGSLVFVSISFIVLMIISSAWLIFYFIQKIRYTNARDRNQRR
 - TIAVGTRMPPKNFSRGSLVFVSISFIVLMIISSAWLIFYFIQKIRYTNARDRNQRR LGDAAKKAISKLTTRT (SEQ ID NO:700); AAKKAISKLTTRTVKKGDKE TDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCPMCKLNIL KALGIVPNLPC (SEQ ID NO:701); TQAVNRRSALGDLAGDNSLGLEPLRTSGI
- 15 SPLPQDGELTPRTGEINIAVTKEWFIIASFGLLSALTLCYMIIRATASLNANEVEW F (SEQ ID NO:702); PLHGVADHLGCDPQTRFFVPPNIKQWIALLQRGNCTF KEKISRAAFHNAVAVVIYNNKSKEEPVTMTHPGTEHIIAVMITELRGKDILSYLE KNISVQMTIAVGTRMPPKNFSRGSLVFVSISFIVLMIISSAWLIFYFIQKIRYTNA RDRNQRRLGDAAKKAISKLTTRTVKKGDKETDPDFDHCAVCIESYKQNDVVRI
- 20 LPCKHVFHKSCVDPWLSEHCTCPMCKLNILKALGIVPNLPCTDNVAFDMERLT RTQAVNRRSALGDLAGDNSLGLEPLRTSGISPLPQDGELTPRTGEINIAVTKEW FIIASFGLLSALTLCYMIIRATASLNANEVEWF(SEQ ID NO:703); and HGVADHLGCDPQTRFFVPPNIKQWIALLQRGNCTFKEKISRAAFHNAVAVVIY NNKSKEE (SEQ ID NO:704). An additional embodiment is the polynucleotide
- fragments encoding these polypeptide fragments. When tested against Jurkat cell lines, supernatants removed from cells containing this gene activated the GAS pathway.

 Thus, it is likely that this gene activates immune cells through the JAKS/STAT signal transduction pathway.

This gene is expressed primarily in macrophage, breast, kidney and to a lesser extent in synovium, hypothalamus and rhabdomyosarcoma.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, schizophrenia and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and neural system, expression of this gene at

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significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to zinc finger protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of schizophrenia, kidney disease and other cancers. The tissue distribution in macrophage, breast, and kidney origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of tumors within these tissues, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 147

The translation product of this gene shares sequence homology with HNP36 protein, an equilibrative nucleoside transporter, which is thought to be important in gene transcription as well as serving as an important component of the nucleoside transport apparatus (See Accession No. 1845345). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

- MSGQGLAGFFASVAMICAIASGSELSESAFGYFITACAVIILTIICYLGLPRLEFYR
 YYQQLKLEGPGEQETKLDLISKGEEPRAGKEESGVSVSNSQPTNESHSIKAILK
 NISVLAFSVCFIFTITIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLG
 RSLTAVFMWPGKDSRWLPSWXLARLVFVPLLLLCNIKPRRYLTVVFEHDAWFI
 FFMAAFAFSNGYLASLCMCFGPKKVKPAEAETAEPSWPSSCVWVWHWGLFS
- 30 PSCSGQLCDKGWTEGLPASLPVCLLPLPSARGDPEWSGGFFF (SEQ ID NO:705); MSGQGLAGFFASVAMICAIASGSELSESAFGYFTTACAVIILTIIC YLGLPRLEFYRYYQQLKLE GPGEQETKLDLISKGEEPRAGKEESGVSVSNSQ PTNESHSI (SEQ ID NO:706); SGVSVSNSQPTNESHSIKAILKNISVLAFSVCFI FTITIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLGRS (SEQ ID
- 35 NO:707),TIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLGRSLTAVF MWPGKDSRWLPSWXLARLVFVPLLLLCNIK PRRYLTVVFEHDA (SEQ ID NO:708); FGPKKVKPAEAETAEPSWPSSCVWVWHWGLFSPSCSGOLCDK

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GWTEGLPASLPVCLLPLPSARGDPEWSGGFFF (SEQ ID NO:709). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in eosinophils and aortic endothelium and to a lesser extent in umbilical vein endothelial cell and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to HNP36 protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of blood neoplasias and other hematopoietic disease.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 148

This gene is expressed primarily in breast cancer cell lines, thymus stromal cells, and ovary.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, endocrine and female reproductive system diseases including breast cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of endocrine disorders. In addition, the tissue distribution in tumors of thymus, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 149

This gene is expressed primarily in retina and ovary and to a lesser extent in brreast cancer cell, epididymus and osteosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal growth disorders, cancer and reproductive system disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neural and reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 382 as residues: Met-1 to Gly-7.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis or treatment of reproductive system disease and cancers.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 150

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One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

MIKDKGRARTALTSSQPAHLCPENPLLHLKAAVKEKKRNKKKKTIGSPKRIQS PLNNKLLNSPAKTLPGACGSPQKLIDGFLKHEGPPAEKPLEELSASTSGVPGLS SLQSDPAGCVRPPAPNLAGAVEFNDVKTLLREWITTISDPMEEDILQVVKYCTD LIEEKDLEKLDLVIKYMKRLMQQSVESVWNMAFDFILDNVQVVLQQTYGSTLK VT (SEQ ID NO:713); MIKDKGRARTALTSSQPAHLCPENPLLHLKAAVKE KKRNKKKKTIGSPKRIQ (SEQ ID NO:714); KRIQSPLNNKLLNSPAKT LPGACGSPQKLIDGFLKHEGPPAEKPLEELSASTSGVPGLSSLQSDPAGCVRPP APNLAGAVEFNDVKTLLREWITTISDPM (SEQ ID NO:715);

TISDPMEEDILQVVKYCTDLIEEKDLEKLDLVIKYMKRLMQQSVE
SVWNMAFDFILDNVQVVLQQTYGSTLKVT (SEQ ID NO:716). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in 12 week embryo and to a lesser extent in hemangiopericytoma and frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, growth disorders and hemangiopericytoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circular and neural system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 383 as residues: Leu-4 to Lys-11.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of growth disorders, hemangiopericytoma and other soft tissue tumors.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 151

The translation product of this gene has been found to have homology to a human DNA mismatch repair protein PMS3. Preferred polypeptide fragments comprise the following amino acid sequence: FCHDCKFPEASPAMNCEP (SEQ ID NO:717). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. R95250).

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lymphoma, immunodeficiency diseases, and cancers resulting from genetic instability. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 384 as residues: Met-1 to Lys-6.

The tissue distribution in neutrophils and the sequence homology indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of Hodgkin's lymphoma, since the elevated expression and secretion by the tumor mass may be indicative of tumors of this type. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. Furthermore, its homology to a known DNA repair protein would suggest gene may be useful in establishing cancer predisposition and prevention in gene therapy applications.

FEATURES OF PROTEIN ENCODED BY GENE NO: 152

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious diseases and lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of inflammation and infectious diseases.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 153

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One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

MASSVPAGGHTRAGGIFLIGKLDLEASLFKSFQWLPFVLRKKC
NFFCWDSSAHSLPLHPLSASCSAPACHASDTHLLYPSTRALCPSIFAWLVAPHS
VFRTNAPGPTPSSQSSPVFPVFPVSFMALIVCXLVCC (SEQ ID NO:720);
MASSVPAGGHTRAGGIFLIGKLDLEASLFKSFQWLPFVLRKKCNFFCWDSSAH
SLPLHPLSASCSAPACHA (SEQ ID NO:721);FAWLVAPHSVFRTNAPGPTPS
SQSSPVFPVFPVSFMALIVCXLVCC (SEQ ID NO:722). An additional embodiment
is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and infectious disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred

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epitopes include those comprising a sequence shown in SEQ ID NO: 386 as residues: Ser-11 to Pro-17.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of infectious diseases and inflammation.

FEATURES OF PROTEIN ENCODED BY GENE NO: 154

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This gene is expressed in multiple tissues including ovary, uterus, adipose tissue, brain, and the liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, uterine, ovarian, brain, and liver cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the female reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnostic or therapeutic uses in the treatment of the female reproductive system, obesity, and liver disorders, particularly cancer in the above tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 155

This gene maps to chromosome 3, and therefore, may be used as a marker in linkage analysis for chromosome 3 (See Accession No. D87452).

This gene is expressed in multiple tissues including brain, aortic endothelial cells, smooth muscle, pituitary, testis, melancytes, spleen, nertrophils, and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological disorders including immunodeficiencies, cancers of the brain and the female reproductive system, as well as cardiovascular disorders, such as

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atherosclerosis and stroke. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution suggest that polynucleotides and polypeptides corresponding to this gene are useful in treatment/detection of disorders in the nervous system, including schizophrenia, neurodegeneration, neoplasia, brain cancer as well as cardiovascular and female reproductive disorders including cancer within the above tissues.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 156

The translation product of this gene shares sequence homology with the human gene encoding cytochrome b561 (See Accession No. P10897). Cytochrome b561 is a transmembrane electron transport protein that is specific to a subset of secretory vesicles containing catecholamines and amidated peptides. This protein is thought to supply reducing equivalents to the intravesicular enzymes dopamine-beta-hydroxylase and alpha-peptide amidase. Preferred polypeptides of the invention comprise the amino acid sequence:

MAMEGYWRFLALLGSALLVGFLSVIFALVWVLHYREGLGWDGSALEFNWHP VLMVTGFVFIQGIAIIVYRLPWTWKCSKLLMKSIHAGLNAVAAILAIISVVAVFE NHNVNNIANMYSLHSWVGLIAVICYLLQLLSGFSVFLLPWAPLSLRAFLMPIHV YSGIVIFGTVIATALMGLTEKLIFSLRDPAYSTFPPEGVFVNTLGLLILVFGALIF WIVTRPQWKRPKEPNSTILHPNGGTEQGARGSMPAYSGNNMDKSDSEL NSEVAARKRNLALDEAGQRSTM (SEQ ID NO:724); as well as antigenic fragments of at least 20 amino acids of this gene and/or biologically active fragments. Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in anergic T-cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune system and metabolism related diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological

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probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product or RNA of this gene is useful for treatment or diagnosis of immune system and metabolic diseases or conditions including Tay-Sachs disease, phenylketonuria, galactosemia, various porphyrias, and Hurler's syndrome.

FEATURES OF PROTEIN ENCODED BY GENE NO: 157

The translation product of this gene shares sequence homology with collagen which is important in mammalian development. This gene also shows sequence homology with bcl-2. (See Accession No. P80988.) Preferred polypeptide fragments comprise the amino acid sequence: PGRAGPSPGLSLQLPAEPGHPAGNLAPL TSRPQPLCRIPAVPG (SEQ ID NO:725). Also preferred are polynucleotide sequences encoding this polypeptide fragment.

This gene is expressed primarily in HL-60 tissue culture cells and to a lesser extent in liver, breast, and uterus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological diseases, hereditary disorders involving the MHC class of immune molecules, as well as developmental disorders and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and reproductive system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those

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comprising a sequence shown in SEQ ID NO: 390 as residues: Ser-39 to Gly-46, Leu-49 to Ala-62.

The tissue distribution and homology to collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hereditary MHC disorders and particularly autoimmune disorders including rheumatoid arthritis, lupus, scleroderma, and dermatomyositis, as well as many reproductive disorders, including cancer of the uterus, and breast tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 158

This gene is expressed primarily in the amygdala region of the brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, a variety of brain disorders, particularly those effecting mood and personality. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and/or diagnosis of a variety of brain disorders, particularly bipolar disorder, unipolar depression, and dementia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 159

This gene is expressed in a variety of tissues and cell types including brain, smooth muscle, kidney, salivary gland and T-cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers of a variety of organs including brain, smooth muscle, kidney, salivary gland and T-cells and cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

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of the above tissues or cells, particularly of the central nervous, urinary, salivary, digestive, and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in brain, smooth muscle, and T-cells indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of various neurological, and cardiovascular disorders, but not limited to cancer within the above tissues. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 160

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The translation product of this gene shares sequence homology with collagen which is thought to be important in cellular interactions, extracellular matrix formation, and has been found to be an identifying determinant in autoimmune disorders.

Moreover, this gene shows sequence homology with the yeast protein, Sls1p, an endoplasmic reticulum component, involved in the protein translocation process in Yeast Yarrowia lipolytica. (See Accession No. 1052828; see also J. Biol. Chem. 271, 11668-11675 (1996).) With mouse, this same region shows sequence homology with the heavy chain of kinesin. (See Accession No. 2062607.) Recently, suppression of the heavy chain of kinesin was shown to inhibits insulin secretion from primary cultures of mouse beta-cells. (See Endocrinology 138 (5), 1979-1987 (1997).) Moreover, kinesin was found associated with drug resistance and cell immortalization. (See 468355.)

Thus, it is likely that this gene also act as a genetic suppressor elements.

This gene is expressed primarily in the greater omentum and to a lesser extent in a variety of organs and cell types including gall bladder, stromal bone marrow cells, lymph node, liver, testes, pituitary, and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the endocrine, gastrointestinal, and immunological systems, including autoimmune disorders and cancers in a variety of organs and cell types.

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Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 393 as residues: Asn-27 to Leu-47, Gln-81 to Lys-88, Asp-93 to Lys-102, Asn-107 to Leu-116, Met-129 to Glu-141, Glu-150 to Asp-157, Lys-176 to Glu-185, Glu-333 to Tyr-349, Cys-393 to Leu-403, Gln-423 to Gly-429.

The tissue distribution in within various endocrine and immunological tissues combined with the sequence homology to a conserved collagen motif indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis of various autoimmune disorders including, but not limited to, rheumatoid arthritis, lupus erthyematosus, scleroderma, dermatomyositis Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 161

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This gene has homology to the tissue inhibitor of metalloproteinase 2. Such inhibitors are vital to proper regulation of metalloproteins such as collagenases (See Accession No. P16368). In addition, this gene maps to chromosome 17, and therefore, may be used as a marker in linkage analysis for chromosome 17 (See Accession No. P16368).

This gene is expressed primarily in several types of cancer including osteoclastoma, chondrosarcoma, and rhabdomyosarcoma and to a lesser extent in several non-malignant tissues including synovium, amygdala, testes, placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, various types of cancer, particularly cancers of bone and cartilage, as well as various autoimmune disorders. Similarly, polypeptides and antibodies directed

to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the musculoskeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in various cancers and the sequence homology to a collagenase inhibitor indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection of various autoimmune disorders such as rheumatoid arthritis, lupus, scleroderma, and dermatomyositis. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 162

This gene is homologous to the mitochondrial ATP6 gene and therefore is likely a homolog of this gene family (See Accession No. X76197).

This gene is expressed primarily in brain tissue.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, a variety of brain disorders, including Down's syndrome, depression, Schizophrenia, and epilepsy. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in brain tissue indicates this gene is useful for diagnosis of various neurological disorders including, but not limited to, brain cancer.

Additionally the gene product may be used as a target in the immunotherapy of cancer in

the brain as well as for the diagnosis of metabolic disorders such as obesity Tay-Sachs disease, phenylketonuria and Hurler's Syndrome.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 163

This gene is expressed primarily in placenta, neutrophils, and microvascular endothelial cells and to a lesser extent in multiple tissues including brain, prostate, spleen, thymus, and bone.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neutropenea and other diseases of the immune system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in placenta indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis various female reproductive disorders. Additionally the gene product may be used as a target in the immunotherapy of various cancers. Because the gene is expressed in some cells of lymphoid and endocrine origin, the natural gene product may be involved in immune functions and metabolism regulation, respectively. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 164

This gene is expressed primarily in neutrophils, monocytes, bone marrow, and 30 fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune system disorders including, but not limited to, autoimmune disorders such as lupus, and immunodeficiency disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

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of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in various immune system tissue indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis of various immunological disorders such as Hodgkin's lymphoma, arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 165

The translation product of this gene shares sequence homology with dystrophin which is thought to be defective in both Duchene and Becker Muscular Dystrophy. 15 Preferred polypeptide fragments comprise the following amino acid sequence: MKLLGECSSSIDSVKRLEHKLKEEEESLPGFVNLHSTETQTAGVIDRWELLQAQ ALSKELRMKQNLQKWQQFNSDLNSIWAWLGDTEEELEQLQRLELSTDIQTIELO IKKLKELQKAVDHRKAIILSINLCSPEFTQADSKESRDLQDRLXQMNGRWDRV CSLLEEWRGLLQDALMQCQGFHEMSHGLLLMLENIDRRKNEIVPIDSNLDAEIL QDHHKQLMQIKHELLESQLRVASLQDMSCQLLVNAEGTDCLEAKEKVHVIGNR 20 LKLLLKEVSRHIKELEKLLDVSSSQQDLSSWSSADELDTSGSVSPXSGRSTPNR QKTPRGKCSLSQPGPSVSSPHSRSTKGGSDSSLSEPXPGRSGRGFLFRVLRAA LPLQLLLLLIGLACLVPMSEEDYSCALSNNFARSFHPMLRYTNGPPPL (SEQ ID NO:726); MKLLGECSSSIDSVKRLEHKLKEEEESLPGFVNLHSTETQTAGVIDR 25 WELLQAQALSKELRMKQNLQKWQQFNSDLNSIWAWLGDTEEELEQLQRLELS TDIQTIELQIK (SEQ ID NO:727); KLKELQKAVDHRKAIILSINLCSPEFTQADSK ESRDLQDRLXQMNGRWDRVCSLLEEWRGLLQDALMQCQGFHEMSHGLLLML ENIDRRKNEIVPIDSNLDAEILQDHHKQLMQIKHELLESQLRVASLQDMSCQL (SEQ ID NO:728); QDMSCQLLVNAEGTDCLEAKEKVHVIGNRLKLLLKEVS RHIKELEKLLDVSSSQQDLSSWSSADELDTSGSVSPXSGRSTPNRQKTPRGKCS 30 LSQPGPSVSSPHS (SEQ ID NO:729); DSSLSEPXPGRSGRGFLFRVLRAAL PLQLLLLLIGLACLVPMSEEDYSCALSNNFARSFHPMLRYTNGPPPL (SEQ ID NO:730). Also preferred are polynucleotide fragments encoding these polypeptide fragments. Furthermore, this gene maps to chromosome 6, and therefore, may be used as a marker in linkage analysis for chromosome 6 (See Accession No. N62896).

This gene is expressed in numerous tissues including the heart, kidney, and brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, musculoskeletal disorders including Muscular Dystrophy and cardiovascular diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the muscle tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to dystrophin indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of Muscular Dystrophy and other muscle disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 166

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This gene is expressed primarily in human cerebellum.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the central nervous system, including Alzheimer's Disease, Parkinson's Disease, ALS, and mental illnesses. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 399 as residues: Pro-20 to Gly-26, Leu-37 to Pro-42, His-57 to Gly-63.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of diseases of the central nervous system and may protect or

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enhance survival of neuronal cells by slowing progression of neurodegenerative diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 167

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Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MKLLICGNYLAPSHSESSRRCCLLCFYPLCLEINFGMKVFLSMPFLVLFQ SLIQED (SEQ ID NO:731). Polynucleotides encoding such polypeptides are also provided. This gene is believed to reside on chromosome 15. Therefore polynucleotides derived from this gene are useful in linkage analysis as chromosome 15 markers.

This gene is expressed primarily in human testes tumor and to a lesser extent in normal human testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the testes, particularly cancer, and other reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of testicular diseases including cancers.

FEATURES OF PROTEIN ENCODED BY GENE NO: 168

This gene is expressed primarily in fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, conditions affecting hematopoietic development and metabolic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the

hepatic system, and fetal hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 401 as residues: His-7 to Trp-17, Leu-19 to Lys-27, Pro-33 to Gly-44, Lys-68 to Gly-74, Lys-85 to Cys-95.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of diseases of the developing liver and hematopoietic system, and act as a growth differentiation factor for hematopoietic stem cells.

FEATURES OF PROTEIN ENCODED BY GENE NO: 169

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The polypeptide encoded by this gene is believed to be a membrane bound receptor. The extracellular domain of which is expected to consist of the following amino acid sequence:

RILLVKYSANEENKYDYLPTTVNVCSELVKLVFCVLVSFCVIKKDHQSRNLKY ASWKEFSDFMKWSIPAFLYFLDNLIVFYVLSYLQPAMAVIFSNFSIITTALLFRIV LKXRLNWIQWASLLTLFLSIVALTAGTKTLQHNLAGRGFHHDAFFSPSNSCLL FRNECPRKDNCTAKEWTFPEAKWNTTARVFSHIRLGMGHVLIIVQCFISSMANI YNEKILKEGNQLTEXIFIQNSKLYFFGILFNGLTLGLQRSNRDQIKNCGFFYGH S (SEQ ID NO:732). Thus, preferred polypeptides encoded by this gene comprise the extracellular domain as shown above. It will be recognized, however, that deletions of either end of the extracellular domain up to the first cysteine from the N-terminus and the first cysteine of the C-terminus, is expected to retain the biological functions of the full-length extracellular domain because the cysteines are thought to be responsible for providing secondary structure to the molecule. Thus, deletions of one or more amino acids from either end (or both ends) of the extracellular domain are contemplated. Of course, further deletions including the cysteines are also contemplated as useful as such polypeptides is expected to have immunological properties such as the ability to evoke

provided.

This gene is expressed primarily in human osteoclastoma and to a lesser extent in hippocampus and chondrosarcoma.

and immune response. Polynucleotides encoding all of the foregoing polypeptides are

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, cancers, particularly those of the bone and connective tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 402 as residues: Met-1 to Cys-6, Ala-41 to Tyr-49, Lys-76 to Lys-84.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis of cancers of the bone and connective tissues, and may act as growth factors for cells involved in bone or connective tissue growth.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 170

Preferred polypeptides encoded by this gene comprising the following amino acid sequence:

NSVPNLQTLAVLTEAIGPEPAIPRXPREPPVATSTPATPSAGPQPLPTGTV LVPGGPAPPCLGEAWALLLPPCRPSLTSCFWSPRPSPWKETGV (SEQ ID NO:733). Polynucleotides encoding such polypeptides are also provided herein.

This gene is expressed primarily in hematopoietic progenitor cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the blood including cancer and autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the blood/circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 403 as residues: Gln-4 to His-10, Pro-25 to His-32.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis of diseases involving growth differentiation of hematopoietic cells.

FEATURES OF PROTEIN ENCODED BY GENE NO: 171

Preferred polypeptides encoded by this gene comprise the following amino acid sequences: ALQLAFYPDAVEEWLEENVHPSLQRLQXLLQDLSEVSAPP (SEQ ID NO:734); and/or CHPPALAGTLLRTPEGRAHARGLLLEAGGA (SEQ ID NO:735). Polynucleotides encoding such polypeptides are also provided. The protein product of this gene shares sequence homology with metallothionines. Thus, polypeptide encoded by this gene are expected to have metallothionine activity, such activities are known in the art and described elsewhere herein.

This gene is expressed primarily in kidney cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the kidney including cancer and renal dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 404 as residues: Ser-47 to Gln-52.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the kidney including kidney failure.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 172

This gene is expressed primarily in 12 week old early stage human.

Therefore, polynocleotides and polynoptides of the invention are used.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for

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differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing embryo, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 405 as residues: Gln-31 to Thr-43, Gly-51 to Ser-58, Pro-65 to Pro-72.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of developmental problems with fetal tissue. The gene may be involved in vital organ development in the early stage, especially hematopoiesis, cardiovascular system, and neural development.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 173

The translation product of this gene shares sequence homology with TGN38, an integral membrane protein previously shown to be predominantly localized to the trans-Golgi network (TGN) of cells.

This gene is expressed primarily in developing embryo and to a lesser extent in cancer tissues including lymphoma, endometrial, protate and colon.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing fetus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 406 as residues: His-65 to Ser-72, Pro-82 to Gly-91, Pro-98 to Glu-118, Ser-126 to Gly-166, Pro-180 to Asp-188, Tyr-209 to Lys-214, Gln-220 to Leu-228.

The tissue distribution and homology to an integral membrane protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for

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diagnosis of cancers and developmental abnormalities where aberrant expression relates to an abnormality.

FEATURES OF PROTEIN ENCODED BY GENE NO: 174

The translation product of this gene shares sequence homology with a dnaJ heat shock protein from E. coli which is allelic to sec63, a gene that affects transit of nascent secretory proteins across the endoplasmic reticulum in yeast.

This gene is expressed primarily in Hodgkin's lymphoma and to a lesser extent in testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 407 as residues: Thr-13 to Trp-21, Arg-74 to Asp-81.

The tissue distribution and homology to dnaJ indicates that polynucleotides and polypeptides corresponding to this gene are useful as a diagnostic for cancer including Hodgkin's lymphoma.

FEATURES OF PROTEIN ENCODED BY GENE NO: 175

This gene is expressed primarily in endothelial cells and to a lesser extent in bone marrow stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases involving angiogenic abnormalities including diabetic retinopathy, macular degeneration, and other diseases including arteriosclerosis and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for treating diseases where an increase or decrease in angiogenesis is indicated and as a factor in the wound healing process.

FEATURES OF PROTEIN ENCODED BY GENE NO: 176

The translation product of this gene shares sequence homology with MAT8 (mouse) which is thought to be important in regulating chloride conductance in cells (particularly in the breast) by modulating the response mediated by cAMP and protein kinase C to extracellular signals.

This gene is expressed primarily in amniotic cells and hematopoeitic cells including macrophages, Neutrophils, T cells, TNF induced aortic endothelium and to a lesser extent in testes, TNF induced epithelial cells, and smooth muscle.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory responses mediated by T cells, macrophages, and/or neutrophils particularly those involving TNF, and also cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEO ID NO: 409 as residues: Thr-19 to Ala-33, Leu-54 to Asp-82, Pro-89 to Ala-97, Pro-100 to Lys-125, Ser-127 to Phe-135, Gly-164 to Leu-169, Cys-173 to Arg-178.

The tissue distribution and homology to mat-8 indicates that polynucleotides and polypeptides corresponding to this gene are useful for modifying inflammatory

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responses to cytokines such as TNF and thus modifying the duration and/or severity of inflammation. Polynucleotides and polypeptides derived from this gene are thought to be useful in the diagnosis and treatment of cancer.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 177

This gene is expressed primarily in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, vascular restenosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating diseases associated with vascular response to injury such as vascular restenosis following angioplasty..

FEATURES OF PROTEIN ENCODED BY GENE NO: 178

One embodiment of the claimed invention comprises:

25 MRPDWKAGAGPGGPPQKPAPSSQRKPPARPSAAAAAIAVAAAEERRLRQRN
RLRLEEDKPAVERCLEELVFGDVENDEDALLRRLRGPRVQEHEDSGDSEVENEA
KGNFPPQKKPVWVDEEDEDEEMVDMMNNRFRKDMMKNASESKLSKDNLKK
RLKEEFQHAMGGVPAWAETTKRKTSSDDESEEDEDDLLQRTGNFISTSTSLPRG
ILKMKNCQHANAERPTVARISICAVPSRCTDCDGCWD (SEQ ID NO:737); or
30 CLEELVFGDVENDEDALLRRLRGPRVQEHEDSGDSEVENEAKGNFPPQKKPV
WVDEEDEDEEMVDMMNNRFRKDMMKNASESKLSKDNLKKRLKEEFQHAMG
GVPAWAETTKRKTSSDDESEEDEDDLLQRTGNFISTSTSLPRGILKMKNCQHA
NAERPTVARISICAVPSRCTDCDGC (SEQ ID NO: 738). LKEKIVRSFEVSPDGS
FLLINGIAGYLHLLAMKTKELIGSMKINGRVAASTFSSDSKKVYASSGDGEVYV
35 WDVNSRKCLNRFVDEGSLYGLSIATSRNGQYVACGSNCGVVNIYNQDSCLQE

TNPKPIKAIMNLVTGVTSLTFNPTTEILAIASEKMKEAVRLVHLPSCTVFSNFPVI KNKNISHVHTMDFSPRSGYFALGNEKGKALMYRLHHYSDF (SEQ ID NO:739);

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and/or KINGRVAASTFSSDSKKVYASSGDGEVYVWDVNSRKCLNRFVDEGSL YGLSIATSRNGQYVACGSNCGVVNIYNQDSCLQETNPKPIKAIMNLVTGVTSLT FNPTTEILAIASEKMKEAVRLVHLPSCTVFSNFPVIKNKNISHVHTMDFSPRSG YFALGNEKGKAL (SEQ ID NO:740).

This gene is expressed primarily in epidydimus and endometrial tumors and to a lesser extent in T cell lymphoma and cell lines derived from colon cancer.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of the reproductive organs including testis and endometrial cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 411 as residues: Ser-67 to Lys-72, Val-87 to Leu-93, Tyr-128 to Pro-141, Asp-204 to Gly-210.

The tissue distribution indicates that the protein products of this gene are useful for treating tumors of the endometrium or epithelial tumors of the reproductive system.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 179

Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MRILQLILLALATGLVGGETRIIKGFECKLHSQPWQAALFEKTRLLCGATLIAPR WLLTAAHCLKPRYIVHLGQHNLQKEEGCEQTRTATESFPHPGFNNSLPNKDH RNDIMLVKMASPVSITWAVRPLTLSSRCVTAGTSCSFPAGAARPDPSYACLTPC DAPTSPSLSTRSVRTPTPATSQTPWCVPACRKGARTPARVTPGALWSVTSLFKA LSPGARIRVRSPESLVSTRKSANMWTGSRRR (SEQ ID NO:741); ETRIIKGFEC KLHSQPWQAALFEKTRLLCGATLIAPRWLLTAAHCLKPRYIVHLGQHNLQKEE GCEQTRTATESFPHPGFNNSLPNKDHRNDIMLVKMASPVSITWAVRPLTLSSR CVTAGTSCSFPAGAARPDPSYACLTPCDAPTSPSLSTRSVRTPTPATSQTPWCVP

ACRKGARTPARVTPGALWSVTSLFKALSPGARIRVRSPESLVSTRKSANMWTG

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SRRR (SEQ ID NO:742); or CKLHSQPWQAALFEKTRLLCGATLIAPRWLLT AAHCLKPRYIVHLGQHNLQKEEGCEQTRTATESFPHPGFNS

(SEQ ID NO:743). The translation product of this gene shares sequence homology with neuropsin a novel serine protease which is thought to be important in modulating extracellular signaling pathways in the brain. Owing to the structural similarity to other serine proteases the protein products of this gene are expected to have serine protease activity which may be assayed by methods known in the art and described elsewhere herein.

This gene is expressed primarily in endometrial tumor and to a lesser extent in colon cancer, benign hypertrophic prostate, and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers of the endometrium or colon and benign hypertrophy of the prostate. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the urogenital or reproductive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 412 as residues: Gly-12 to Ser-22, Pro-34 to Ser-53.

The tissue distribution and homology to serine proteases indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing or treating hyperpoliferative disorders such as cancer of the endometrium or colon and hyperplasia of the prostate.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 180

Preferred polypeptide encoded by this gene comprise the following amino acid sequence: VLQGRYFSPILEMRRLRPEGXXNLPGGSRAQKEPRQDLTLVLWPHC PHFAMTRSYVPTKQCMVQGSFYCIFIFKGPVQNWC (SEQ ID NO:744).

35 Polynucleotides encoding such polypeptide are also provided.

This gene is expressed primarily in fetal brain

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, identifying and expanding stem cells in the CNS. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for detecting and expanding stem cell populations in the (or of the) central nervous system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 181

This gene is expressed primarily in early stage human brain and a stromal cell line.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities of the CNS. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 414 as residues: Gln-42 to Gln-47, Gln-54 to Pro-60.

The tissue distribution indicates that the protein products of this gene play a role in the development of the central nervous system. Therefore this gene and its products

are useful for diagnosing or treating developmental abnormalities of the central nervous system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 182

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Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MPIIDQVNPELHDFMQSAEVGTIFALSWLITWFGHVLSDFRHVVRLYDF FLACHPLMPIYFAAVIVLYREQEVLDCDCDMASVHHLLSQIPQDLPYETLISRXE TFLFSFPHPNLLGRPLPNSKLRGRQPLLSKTLSWHQPSRGLIWCCGSGXRGLL RPEDRTKDVLTKPRTNRFVKLAVMGLTVALGAAALAVVKSALEWAPKFQLQL FP (SEQ ID NO:745); or CPEFFIPATLPCPFVFAFTSEASSRAYLTQRGPGGLAQ NLMPLPVGFWMGSLPPPWCWRKWVSEACSCFC (SEQ ID NO:746) These polypeptides are structurally similar to various TGF-beta family members. Thus, this polypeptide is expected to have a variety of activities in the modulation of cell growth and proliferation.

This gene is expressed primarily in osteoclastoma, microvascular endothelium, and bone marrow derived cell lines.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematological diseases particularly involving aberrant proliferation of stem cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 415 as residues: Ser-33 to Ala-39.

The tissue distribution indicates that the protein products of this gene is useful for treating disorders of the progenitors of the immune system. Applications include in vivo expansion of progenitor cells, ex vivo expansion of progenitor cells, or the treatment of tumors of the circulatory system, such as lymphomas.

FEATURES OF PROTEIN ENCODED BY GENE NO: 183

This gene maps to chromosome 17 and therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 17. In specific embodiments, polypeptides of the invention comprise the sequence:

- 10 GVFEVPKQNGKYETGQLFLHSIFGYRGVVLFPWQARLXDRDVASAAPEKAEN PAGHGSKEVKGKTHTYYQVLIDARDCPHISQRSQTEAVTFLANHDDSRALYAIP GLDYVSHEDILPYTSTDQVPIQHELFERFLLYDQTKAPPFVARETLRAWQEKNH PWLELSDVHRETTENIRVTVIPFYMGMREAQNSHVYWWRYCIRLENLDSDVVQ LRERHWRIFSLSGTLETVRGRGVVGREPVLSKEQPAFQYSSHVSLQASSGHMW
- 15 GTFRFERPDGSHFDVRIPPFSLESNKDEKTPPSGLHW (SEQ ID NO:751);
 MAACTARRPGRGQPLVVPVADXGPVAKAALCAA (SEQ ID NO:752);
 VLETVGVFEVPKQNGKYETGQLFLHSIFGYRGVVL (SEQ ID NO:757);
 GLDYVSHEDILPYTST (SEQ ID NO:758); DVHRETTENIRVTVIPFYM (SEQ ID NO:759); WWRYCIRLENLDSDVVQLRER (SEQ ID NO:760); and/or PAFQYSS
- 20 HVSLQASSGHMWGTFRFER (SEQ ID NO:761). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in gall bladder, prostate, and fetal brain, and to a lesser extent in a few tumor and fetal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as 25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, growth related disorders such as cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate, gall bladder, and fetal brain, 30 expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., 35 the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of growth-related disorders, such cancers.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 184

In specific embodiments, polypeptides of the invention comprise the sequence:SLCCPEGAEGC (SEQ ID NO:762) and/or QLKKTHYDRPCP (SEQ ID NO:763). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in stromal cell, tonsil, and glioblastoma and to a lesser extent in some other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune and inflammatory disorders and glioblastoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the stromal cells, tonsil, and glioblastoma expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Additionally, it is believed that the product of this gene regulates pancreatic cell differentiation into beta cells. Accordingly, polynucleotides and polypeptides of the invention are useful in the treatment of insulindependent diabetes mellitus and associated conditions e.g. pancreatic hypofunction and the prevention, as well as the treatment of undifferentiated type pancreatic cancers. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 417 as residues: Pro-27 to Ala-32.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune and inflammatory disorders and glioblastoma.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 185

This gene is expressed primarily in hepatocellular carcinoma and to a lesser extent in other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 418 as residues: Gly-32 to Lys-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of liver diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 186

This gene is expressed primarily in hippocampus and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neutronal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hippocampus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal disorders.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 187

This gene is expressed primarily in bone cancer and hippocampus and to a lesser extent in osteoclastoma and other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, bone-related disorders and neuronal diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the bone, ostoeclast, and hippocampus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of bone-related disorders and neuronal diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 188

This gene maps to chromosome 4 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 4.

This gene is expressed primarily in neuronal tissues such as hippocampus, spinal cord, and hypothalamus and to a lesser extent in a few other tissues such as ovary.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 189

This gene maps to chromosome 10, therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 10.

This gene is expressed primarily in neuronal tissues and immune tissues, and to a lesser extent in a few other tissues such as skin tumor, lung etc.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal and immune-related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal and immune-related tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 422 as residues: Pro-19 to Asp-25.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal and immune-related disorders.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 190

The translation product of this gene shares sequence homology with human N33, a gene located in a homozygously deleted region of human metastatic prostate cancer which is thought to be important in prevention of prostate cancer. In specific embodiments, polypeptides of the invention comprise the sequence:

- 30 AQRKKEMVLSEKVSQLMEWTNKRPVIRMNGDKFRRLVKAPPRNYSVIVMFTA LQLHRQCVVCKQADEEFQILANSWRYSSAFTNRIFFAMVDFDEGSDVFQMLNM NSAPTFINFPAKGKPKRGDTYELQVRGFSAEQIARWIADRTDVNIRVIRPPNMA ARWRFWCVSVT (SEQ ID NO:765); MVVALLIVCDVPSAS (SEQ ID NO:766); AQRKKEMVLSEKVSQL (SEQ ID NO:767); MEWTNKRPVIRMNGDKF (SEQ ID:768); RRLVKAPPRNYSVIVMFTALQLHRQCVVCKQADEEFQILANSWRY
- SSAFTNRIFFA (SEQ ID NO:769); MVDFDEGSDVFQMLNMNSAPTFINFPAK
 GKP (SEQ ID NO:770); KRGDTYELQVRGFSAEQIARWIADRTDVNIRVIRPPN

(SEQ ID NO:771); and/or YAGPLMLGLLLAVIGGLVYLRRVIWNFSLIKLDGLLQL CVLCLL (SEQ ID NO:772). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in infant adrenal gland prostate cell line and to a lesser extent in a few other tissues like liver, smooth muscle etc.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate cancer and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate and adrenal gland, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 423 as residues: Pro-34 to Gly-43, Arg-113 to Pro-120.

The tissue distribution and homology to N33 indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for prostate cancer and endocrine disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 191

This gene is expressed primarily in T cell and to a lesser extent in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 424 as residues: Trp-3 to Phe-9.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 192

This gene maps to chromosome 6, therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 6. Neural activity and neurotrophins induce synaptic remodeling in part by altering gene expression. This gene is believed to be a glycosylphoshatidylinositol-anchored protein encoded by a hippocampal gene and to possess neural activity. This molecule is believed to be expressed in postmitotic-differentiating neurons of the developing nervous system and neuronal structures associated with plasticity in the adult. Message of this gene is believed to be induced by neuronal activity and by the activity-regulated neurotrophins BDNF and NT-3. The product of this gene is believed to stimulate neurite outgrowth and arborization in primary embryonic hippocampal and cortical cultures and to act as a downstream effector of activity-induced neurite outgrowth. In specific embodiments, polypeptides of the invention comprise the sequence: DAVFKGFSDCLLKLGDS (SEQ ID NO:773); CQEGAKDMWDKLRKESKNLN (SEQ ID NO:774); VLLVSLSAALATWLSF (SEQ ID NO:775); MGLKLNGRYISLILAVQIAYLVQAVR AAGKCDAVFKGFSDCLLKLGDS (SEQ ID NO:776); PAAWDDKTNIKTVCTYW EDFHSCTVTALTDCQEGAKDMWDKLRKESKNLNIQGSLFELCGSGNGAAGSL LPAFPVLLVSLSAALATWLSF (SEQ ID NO:777); and/or MGLKLNGRYISLILA VOIAYLVQAVRAAGKCDAVFKGFSDCLLKLGDSXXXXXPAAWDDKTNIKTVC TYWEDFHSCTVTALTDCQEGAKDMWDKLRKESKNLNIQGSLFELCGSGNGAA GSLLPAFPVLLVSLSAALATWLSF (SEQ ID NO:778). Polynucleotides encoding this polypeptide are also encompassed by the invention.

This gene is expressed primarily in human placenta, endometrial tumor and tissues of the central nervous system (CNS).

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, relating to reproductive disorders, cancers and neurological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and neurological disorders, expression of this gene at significantly higher

or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 425 as residues: Asp-47 to Asp-63, His-75 to Tyr-80, Pro-83 to Tyr-89.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of reproductive disorders such as endometrial tumors. Expression of this gene in tissues of the CNS and its strong homology to Neuritin suggest that the protein product from this gene may also be used in the treatment and diagnosis of neurological disorders and in the regeneration of neural tissues, e.g., following injury.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 193

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The translation product of this gene shares sequence homology with tenascin which is thought to be important in development. The translation product of this gene is believed to be a ligand of the fibroblast growth factor family. FGF ligand activity is known in the art and can be assayed by methods known in the art and disclosed elsewhere herein.

This gene is expressed primarily in endometrial tumors, and other types of tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 426 as residues: Gly-29 to Glu-34, Arg-71 to Arg-76, Thr-176 to Cys-182, Gly-184 to Glu-199.

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The tissue distribution and homology to tenascin indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancers.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 194

In specific embodiments, polypeptides of the invention comprise the sequence: MNSAAGFSHLDRRERVLKLGESFEKQPRCASTLC (SEQ ID NO:779). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in fetal human lung and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lung development and respiratory disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the respiratory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in fetal lung and neutrophils indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of lung and immunity related diseases, for example, lung cancer, viral, fungal or bacterial infections (e.g. lesions caused by tuberculosis), inflammation (e.g. pneumonia), metabolic lesions etc.

FEATURES OF PROTEIN ENCODED BY GENE NO: 195

This gene is expressed primarily in breast lymph node.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at

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significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immunal disorders.

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 196

This gene maps to chromosome 5 and accordingly, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 5. The translation product of this gene shares sequence homology with human M-phase phosphoprotein 4 which is thought to be important in phosphorylation and signal transduction processes. In 15 specific embodiments, polypeptides of the invention comprise the sequence: TIYPTEEELQAVQKIVSITERALKLVSD (SEQ ID NO:780); RALKGVLRV GVLAKGLLLRGDRNVNLVLLC (SEQ ID NO:781); ALAALRHAKWFQARAN GLQSCVIIIRILRDLCQRVPTWS (SEQ ID NO:782); GDALRRVFECISSGIIL (SEQ ID NO:783); LAFRQIHKVLGMDPLP (SEQ ID NO:784); and/or TIYPTEEELQAVQ 20 KIVSITERALKLVSDSLSEHEKNKNKEGDDKKEGGKDRALKGVLRVGVLAKG LLLRGDRNVNLVLLCSEKPSKTLLSRIAENLPKQLAVISPEKYDIKCAVSEAAIIL NSCVEPKMQVTITLTSPIIREENMREGDVTSGMVKDPPDVLDRQKCLDALAALR HAKWFQARANGLQSCVIIIRILRDLCQRVPTWSDFPSWAMELLVEKAISSASSP QSPGDALRRVFECISSGIILKGSPGLLDPCEKDPFDTLATMTDQQREDITSSAQFA 25 LRLLAFRQIHKVLGMDPLPQMSQRFNIHNNRKRRRDSDGVDGFEAEGKKDKK DYDNF (SEQ ID NO:785). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in Human Hippocampus and to a lesser extent in Prostate, Human Frontal Cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders related to reproductive system and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system and nervous system, expression of this gene at significantly higher or lower

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levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to human M-phase phosphoprotein 4 indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of reproductive and nervous system disorders.

10 FEATURES OF PROTEIN ENCODED BY GENE NO: 197

In specific embodiments, polypeptides of the invention comprise the sequence: MGSQHSAAARPSSCRRKQEDDRDG (SEQ ID NO:786); LLAEREQEEAIAQFPYVEFTGRDSITCLTC (SEQ ID NO:787); and/or QGTGYIPTEQVNELVALIPHSDQRLRPQRTKQYV (SEQ ID NO:788).

Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in Human Primary Breast Cancer and to a lesser extent in Human Adult Spleen, Hodgkin's Lymphoma I, Salivary Gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer and immunal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 430 as residues: Ser-126 to Gly-138.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of cancer and immunal disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 198

This gene is expressed primarily in monocytes.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, blood cell disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of blood cell disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 199

This gene is expressed primarily in Human Ovary and Synovia and to a lesser extent in Human 8 Week Whole Embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive and developmental disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and developmental system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of reproductive and developmental disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 200

This gene maps to chromosome 8 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 8. The translation product of this gene shares limited sequence homology with collagen proline rich domain.

This gene is expressed primarily in CNS.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 433 as residues: Pro-35 to Asp-41.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological diseases.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 201

Translation product of this gene shares homology with a mammalian histone H1a protein. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: ARLNVGRESLKREMLKSQGVKVSESPMGAR HSSWPEGAAFCKKVQGAQMQFPPRR (SEQ ID NO:789); ARLNVGRESLKR EML (SEQ ID NO:790); LKSQGVKVSESPMGARHSSW (SEQ ID NO:791); AFCKKVQGAQMQFPPRR (SEQ ID NO:792). An additional embodiment is the polynucleotide fragments encoding these polypeptide (See Accession No. pirlS24178) fragments.

This gene is expressed primarily in neutrophils.

35 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in vital immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 202

This gene is expressed primarily in neutrophils.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 203

This gene is expressed primarily in Neutrophils.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious disorders, immune disorders, and cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 436 as residues: Thr-31 to Lys-36.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of infectious disorders, immune disorders, and cancers. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

FEATURES OF PROTEIN ENCODED BY GENE NO: 204

This gene maps to chromosome 16 and therefore polynucleotides of the invention can be used in linkage analysis as markers for chromosome 16. The translation product of this gene shares sequence homology with lactate dehydrogenase which is thought to be important in lactate metabolism.

This gene is expressed primarily in human tonsils and to a lesser extent in Spleen, and Neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders, infectious disorders, and cancers. Similarly,

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polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune disorders, infectious disorders, and cancers, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 437 as residues: Gly-7 to Ser-12.

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The tissue distribution and homology to lactate dehydrogenase gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders, infectious disorders, and cancers.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 205

The translation product of this gene shares sequence homology with Gcap1 protein which is developmentally regulated in brain.

This gene is expressed primarily in placenta and endometrial tumor and to a lesser extent in several other tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, vasculogenesis/angiogenesis and tumorigenesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system and tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Gcap1 protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorder or dysfunction of vascular system of tumorigenesis.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 206

In specific embodiments, polypeptides of the invention comprise the sequence MPYAQWLAENDRFEEAQKAFHKAGRQREA (SEQ ID NO:799); VQVLEQLTNNAVAESRFNDAAYYYWMLSMQCLDIAQD (SEQ ID NO:794); PAQKDTMLGKFYHFQRLAELYHGYHAIHRHTEDP (SEQ ID NO: 795); FSVHRPETLFNISRFLLHSLPKDTPSGISKVKILFT (SEQ ID NO:800); LAKQSKALGAYRLARHAYDKLRGLYIP (SEQ ID NO:796); ARFQKSIELG TLTIRAKPFHDSEELVPLCYRCSTNN (SEQ ID NO: 797); and/or PLLNNLGNVC INCRQPFIFSASSYDVLHLVEFYLEEGITDEEAISLIDLEVLRPKRDDRQLEICKQQ LPDSCG (SEQ ID NO:798). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of male reproductive and endocrine disorders.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 207

This gene is expressed in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lung diseases such as cystic fibrosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

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of the above tissues or cells, particularly of the respiratory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 440 as residues: Tyr-49 to Cys-54.

The tissue distribution indicates that polynucleotides and polypeptides

corresponding to this gene are useful for detection and treatment of disorders associated with developing lungs particularly in premature infants where the lungs are the last tissues to develop. The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of lung tumors since the gene may be involved in the regulation of cell division,

particularly since it is expressed in fetal tissue. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

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25	87	121	49	56	218	30		Last AA of ORF

33	32	31	30	30	29	28		Gene No.	
HTWCI46	HTWBY48	HJPCD40	HTSEV09	HTPBW79	HTOAM21	HTGEU09		cDNA Clone ID	
97974 04/04/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	209511 12/03/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	04/04/97 209080 05/29/97	Deposit Nr and Date	ATCC
pSport1	pSport1	Uni-ZAP XR	pBluescript	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
43	42	41	222	40	39	38		ןĦ	NT
1821	1094	704	1404	1515	812	872	l	Total NT Seq.	
892	-	22	-	811				Clone Seq.	of Of
1647	1094	704	1265	1507	812	872		Clone Clone Seq. Seq.	5' NT 3' NT
56	32		92	302	41	74		Say	νī
56	32	117	%	302	41	74		of AA of ID of of of art Signal NO: Sig Sig Sig	5' NT of First
266	265	264	445	263	262	261		≺ö₽,	AA
_		9			1			of Sig Pep	First AA
26	34	18	19	24	30	1∞			Last AA
27	35	19	20	25	31	19			First AA
28	53	127	415	362	43	28		ORF A	Last

39	38	37	. 36	35	35	34		Gene No.
HBMSN25	HATEF60	HAGFB60	HADAE74	HWTBF59	HWTBF59	HTXGI75		cDNA Clone ID
97974	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	209080 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pSport1	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector
49	48	47	46	223	45	44		× Ö. B.Ö. N.J.
1742	2432	840	2421	707	983	1024		Total NT Seq.
1165	1193	1	664	488	779	30		5' NT of Clone Seq.
1742	2246	840	1587	707	983	1024		5' NT 3' NT of Of Clone Clone Seq. Seq.
1742 1165 1742 1207	1491	97	710	514	85			5' NT of Start Codon
1207	1491	97	710	514	85	167		of AA First SEQ AA of ID Signal NO: Pep Y
272	271	270	269	446	268	267		≺ NO: SEO SEO AA
E	1			_		-		First AA of Sig Pep
23	17	30		4	30	20		Last AA of Sig Pep
24	180	<u>y</u>		42	ω	21		First AA of Secreted Portion
31	51	48	2	2	221	25		Last AA of ORF

45	4	43	42	41	40		Gene No.
HCESF40			HMDAN54	HCE3J79	HCDAR68		cDNA Clone ID
97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	04/04/97 209080 05/29/97	ATCC Deposit Nr and Date
pBluescript	Uni-ZAP XR		Vector				
55	54	53	52	51	50		X D D N N N N N N N N N N N N N N N N N
990	948	1558	1856	1328	1487		Total NT Seq.
99	1	310	725	251	181		5' NT of Clone Seq.
990	948	1408	1853	1328	1455		3' NT of Clone Seq.
193	9	393	928	525	325		5' NT of Start Codor
193	9	393	928	525	325		of AA I of SEQ AA of ID Signal NO: Pep Y
278	277	276	275	274	273		Y. NO: SEQ SEQ
	-	į.	1	-	1		First AA of Sig Pep
32	23		33		35		Last AA of Sig Pep
33	24		34		36		First AA of Secreted Portion
256	65	1	50	21	36		Last AA of ORF

51	50	49	48	47	46	45	Gene No.
HCWBB42	HCUDC07	HCRAF32	HCNAP62	HCMSX86	HCFMV39	HCESF40	cDNA Clone ID
97975 04/04/97 209081	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	ATCC Deposit Nr and Date
ZAP Express	ZAP Express	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	pSport1	pBluescript	Vector
61	60	59	58	57	56	224	× N B SE N
618	478	1215	814	1052	1603	1384	Total NT Seq.
-	grapes.	1215 257	1	5	-	99	5' NT of Clone Seq.
618	478	1215	558	786	1296	1384	5' NT 3' NT of of Clone Clone Seq. Seq.
212	147		93	12	96	193	5' NT of Start Codor
212	147	356	93	12	96	193	5' N' of First AA c Signa Pep
284	283	282	281	280	279	447	≺QEQ\$
<u> </u>		-	1	1	1	-	First AA of Sig Sig Pep
35	36	19	22	28	29		Last AA of Sig Pep
36	37	20	23	29	30	33	First AA of Secreted Portion
74	69	20	42	32	102	205	Last AA of ORF

58	57	56	55	54	53	52		Gene No.
		6		+>				ne o.
нЕ9НU17	HE6EU50	HE2OF09	HE2GS36	HE2AY71	HE2AV74	HDTAB05		cDNA Clone ID
97975 04/04/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pCMVSport 2.0		Vector
68	67	66	65	2	63	62		X D S N
2483	1152	1866	774	588	780	751		Total NT Seq.
1577	117	1313	272	21	283	-		5' NT of Clone Seq.
2448	686	1866	774	588	780	751		3' NT of Clone Seq.
1620	237	1596		169		257		5' NT of Start Codon
1620	237	1596	445	169	433	257		of AA I First SEQ AA of ID Signal NO:
291	290	1		287	286	285		≺ö BŠ¥
			_			-		First AA of Sig Pep
	20					21		Last AA of Sig Pep
	21					22		First AA of Secreted Portion
4	34		37	16	16	32		Last AA of ORF

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65	64	63	62	61	60	59		Gene No.
HFVHY45	HFGAB89	HFEBA88	HEMAE80	HELDY74	HEBBW11	HE9ND48		cDNA Clone ID
97975	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	209081 05/29/97	ATCC Deposit Nr and Date
pBluescript	Uni-ZAP XR		Vector					
75	74	73	72	71	70	69		× N E S Y
831	1069	785	996	932	865	536		Total NT Seq.
-	196	464	,		647			5' NT of Clone Seq.
831	1047	785	945	932	865	536		5' NT 3' NT of of Clone Clone Seq. Seq.
	295	356	12	201		83		5' N' of Start
88	295	356	12	201	388 8	83		of AA First land of AA First land of AA AA AA AA Of D Of Signal NO: Signal NO
298	297	296	295	294	293	292		, A SE SE SE SE SE SE SE SE SE SE SE SE SE
	-	-	-	-				First AA of Sig Pep
30	32	29	24	17	30	36		Last AA of Sig Pep
31	33	30	25	-	31	37		First AA of Secreted Portion
76	34	57	136	33	135	43		Last AA of ORF

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71	70	69	89	67	66		Gene No.
HHGCN69	HHFHR32	ннғнл59	HHFCF08	нсвв 269	НСВАЈ93		cDNA Clone II
CN69	1R32	H129	CF08	BQ69	AJ93		cDNA Clone ID
97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	04/04/97 209081 05/29/97	ATCC Deposit Nr and Date
75 4/97 381 9/97	4/97 4/97 081 9/97	775 4/97 081 9/97	775 4/97 081 9/97	775 4/97 081 9/97	775 4/97 081 9/97	4/97 081 9/97	and CC
Lamt	Uni-Z	Uni-2	Uni-2	Uni-2	Uni-2		<
Lambda ZAP II	Uni-ZAP XR		Vector				
4	(R	(R	(R	R	- R		
82	80	79	78	77	76		×öesi
1440	1378	661	1133	1274	590		Total NT Seq.
298			4	,			5' NT of Clone Seq.
1440	1378	661	1042	1273	590		5' NT 3' NT of of Clone Clone Seq. Seq.
532		192	175	105			S' N'I
532	358	192	175	105	233		of AA First Of AA First First SEQ AA AA of ID of Signal NO: Sig Pep Y Pep
304	303	302	301	300	299		≺ö BŠ SŠ SŠ
1	1	-	1		-		First AA of Sig Pep
23		29	23	24	<u>ယ</u> တ		Last AA of Sig Pep
24		30	24	25	39		First AA of Secreted Portion
34	13	112	30	43	94		Last AA of ORF

			_	- 1							
82	18	80	79	78	77	76	75	74	73	72	Gene No.
HNGBT31	HNFJH45	HNFAE54	HMSKS35	HMEJE31	HKMNC43	HKIXL73	HJPAV06	HHSEG23	HHPFD63	HHGDO13	cDNA Clone ID
97976 04/04/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	ATCC Deposit Nr and Date								
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	pBluescript	pBluescript	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	Vector
92	91	90	68	88	87	86	85	84	83	82	X D D S N I
639	575	1533	1102	655	908	1036	684	573	1706	1381	Total NT Seq.
_	1	665	1	1	-	165	199	-	182	766	5' NT of Clone Seq.
639	575	1518	1102	655	908	1036	684	573	1644	1371	5' NT 3' NT of of Clone Clone Seq. Seq.
224	275	347		165	139		323	160	257	993	5' NT of Start Codon
224	275	347	228	165	139	690	323	160	257	993	
315	314	313	312	311	310	309	308	307	306	305	≺ö BÖ SEÖ A
_	_	-	_	-	-	-	-	-		_	First AA of Sig Pep
28	30	26	26	ယ္သ	18	32	27	18	24	23	Last AA of Sig Pep
29	31	27	27	34	19	33	28	19	25	24	First AA of Secreted Portion
8	67	293	49	2	801	114	33	71	<u>∞</u>	34	Last AA of ORF

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91	90	89	88	87	86	85	84	83	Gene No.	
HPCAL49	HPBCU51	HOSDI92	HOSBZ55	HOGAR52	HNHFL57	HNHDW42	HNGJG84	HNGIN60	cDNA Clone ID	
97977 04/04/97 209082	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97976 04/04/97	97976 04/04/97	97976 04/04/97	97976 04/04/97	Nr and Date	ATCC
Uni-ZAP XR	pBluescript SK-	Uni-ZAP XR	Uni-ZAP XR	pCMVSport 2.0	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Vector	
101	100	99	98	97	96	95	94	93	׊E	SEO
784	599	1935	1416	1985	844 4	426	526	744	,	3
1	-	141	69	453		_	-		Seq.	of S' NT
784	599	772	1416	1985	844	426	526	744	Seq.	5' NT 3' NT of of
	86		246	533	98	168	268	225	Start Codor	S' NT
280	8	274	246	533	98	168	268	225	Signal NO:	of First
324	323	322	321	320	319	318	317	316	ĸöŧ	SEQ
_	_	-	-	-					Sig	A First
18	27	20	32	17	25	28	29	1	10 44	Last AA
19	28	21	33	188	26	29	30	44	Secreted Portion	First AA
43	119	58	54	285	01		38	1		Last

	·				· · · · · · · · · · · · · · · · · · ·				
97	96	95	95	94	93	92		Gene No.	
HRGBR28	HRDFB85	HPWAN23	HPWAN23	нрмвQ32	НРНАС83	HPFCR13		cDNA Clone ID	
97977 04/04/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	05/29/97	Deposit Nr and Date	ATCC
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
107	106		105	104	103	102		ןĦ	NT
1167	1705	226 2057	2066	1351	2218	1035		Total NT Seq.	
611	23	-	51	-	840	602		Clone Seq.	of IN 'S
1167	1697	1954	2052	1351	2182	1035		,	5' NT 3' NT
53		1		18	1035	859		of Start Codor	5' NT
53	233	220	270	.	1035	859		AA of ID Signal NO: Pep Y	5' NT of First
330	329	449	328	327	326	325		⊀ö.£	AA SEQ
	_	,	-			1		of Sig Pep	First AA
_	21	29	29	23	17	32		of Sig Pep	Last AA
2	22	30	30	24	18	33		of Secreted Portion	First AA
263	201	315	537	86	17	58		ORF & A	Last

102	101	100	100	99	98	98		Gene No.
HTEFU09	HSXCS62	HSXBT86	HE8EU04	HSPAH56	HSKGN81	HSKGN81		cDNA Clone ID
97977 04/04/97 209082	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209746 04/07/98	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209082 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pSport1	pBluescript	pBluescript		Vector
112	111		110	109	227	108		× NO: SEO NT
2198	2249	228 2143	2632	611	2084	1907		Total NT Seq.
228		53	294		335	151		5' NT of Clone Seq.
2158	1953	1096	2632	576	2084	1432		5' NT 3' NT of of Clone Clone Seq. Seq.
400	90		337	229		353		S' N' of Start
400	90	235	337	229	537	353		of AA First I of AA First I First SEQ AA AA of ID of Signal NO: Sign Pep Y Pep
335	334	451	333	332	450	331		YO. BEO AA
_	-	_	1	1	-	1		First AA of Sig Pep
	18		25	25	19	23		Last AA of Sig Pep
	19		26	26	20	24		First AA of Secreted Portion
23	199	9	333	47	23	260		Last AA of ORF

109	108	107	106	105	104	103		Gene No.
HTSHE40	HTSGM54	HTPCN79	нтоеу16	HTGEW91	HTGEP89	НТЕКМ35		cDNA Clone ID
97977 04/04/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	05/29/97	ATCC Deposit Nr and Date
pBluescript	pBluescript	Uni-ZAP XR		Vector				
119	118	117	116	115	114	113		× N E SEO N T
1101	1133	503	1965	3684	703	1043		Total NT Seq.
118	316	_	127	526	-	40		5' NT of Clone Seq.
956	1069	503	1915	1338	703	1043		5' NT 3' NT of of Clone Clone Seq. Seq.
218			202	584	285	320		5' NT of Start Codor
218	423	1	202	584	285	320		of AA First SEQ AA of ID Signal NO: Pep Y
342	341	340	339	338	337	336		Y. DEQ AA
	-	1	-	1	1	₽		First AA of Sig Pep
31	12	7	27	24	29	20		Last AA of Sig Pep
32	13	∞	28	25	30	21		First AA of Secreted Portion
89	84	70	38	37	94	142		Last AA of ORF

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911	115	14	113	112	11	110		Gene No.	
HE6EL90	HDTAW95	HCEVR60	нсезQ10	HUKFC71	HTWBY29	HTWAF58		cDNA Clone ID	
209007	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209082 05/29/97	ATCC Deposit Nr and Date	
Uni-ZAP XR	pCMVSport 2.0	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	pSport1	Lambda ZAP II		Vector	
126	125	124	123	122		120		× Ö Ð SE	į.
126 1517	1288	124 1390	1542	994	2635	282		Total NT Seq.	
	412	82		ı	1593			of Clone Seq.	יין
1452	1288	1390	1542	932	2635 1593 2489	282		of Clone Seq.	יין יינ
243	571	127	143		1654	137		5' NT of Start Codon	
243	571	127	143	272	1654	137		First SEQ AA of ID Signal NO: Pep Y	5' NT
349	348	347	346	345	344	343		≺Ö∄ÖŞ SÖĞ	>
1	_	1	_	-	1	1		AA AA OF Sig Pep Pep	
		32	25	15	25	25		AA of Sig Pep	1
		33	26	16	26	26		First AA of Secreted Portion	
9	16	153	63	221	55	48		Last AA of ORF	

122	121	120	119	811	117		Gene No.
HLTER03	HIBED17	HHPTD20	HFXBW82	HERAH36	HELBU29		cDNA Clone ID
209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	04/28/97 209083 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Other	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR		Vector
132	131	130	129	128	127		× N B S N
990	1950	472	1275	300	1073		Total NT Seq.
-	284	51		155			5' NT 3' NT of of Clone Clone Seq. Seq.
990	1927	472	1275	300	1073		3' NT of Clone Seq.
78	395		56	202			5' NT of Start Codon
78	395	243	56	202	776		5' NT of First AA of Signal Pep
355	354	353	352	351	350		≺ö ÐSA
-	-	_	_		_		First AA of Sig Pep
22	72		23				Last AA of Sig Pep
23	73		24				First AA of Secreted Portion
34	245	32	6	17	13		Last AA of ORF

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129	128	127	126	125	124	123	Gene No.
H6EAA53	HUKCO64	HSUBW09	HRGBR18	HPWAZ95	НРМСЈ92	HOABL56	cDNA Clone ID
209007 04/28/97 209083	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Vector				
139	138	137	136	135	134	133	× Ö. BÖ SEÖ N. T. V. V. T. V.
643	1777	137 1021	582	323	705	1720	Total NT Seq.
303	439	1	1		28	565	5' NT of Clone Seq.
643	1777	1021	582	323	705	1720	3' NT of Clone Seq.
		153		88	106	660	5' NT of Start Codon
313	521	153	16	88	106	660	of AA First SEQ AA of ID Signal NO: Pep Y
362	361	360	359	358	357	356	YO. DEQ SEQ AA
-	•	 _	-		•	J	First AA of Sig Pep
7		32	17	27	28	18	First Last AA AA of of Sig Sig Pep Pep
∞		33	18	28	29	19	First AA of Secreted Portion
31	2	56	30	78	98	21	Last AA of ORF

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135	134	134	133	132	131	130		Gene No.
HBMTD81	нвссв91	HAIBP89	HALSQ59	HALSK07	HAGAO39	HAGAIII		cDNA Člone ID
209008 04/28/97 209084 05/29/97	209007 04/28/97 209083 05/29/97	unknown 05/18/98	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector
145	229	144	143	142	141	140		XO: SEO NT
1082	229 1025	144 2243	300	1468	721	1220		Total NT Seq.
163	409	173	4	125		-		5' NT of Clone Seq.
1082	1025	2243	300	1468	721	1220		5' NT 3' NT of of Clone Clone Seq. Seq.
357	624	311	101	210				5' NT of Start Codon
357	624	311	101	210	415	127		5' N' of First AA c Signa Pep
368	452	367	366	365	364	363		Y P
-		1	1	_	•			First AA of Sig Pep
	20	27	22	29		16		Last AA of Sig Pep
	21	28	23	30		17		First AA of Secreted Portion
30	25	317	66	33	14	27		Last AA of ORF

							
142	141	140	139	138	137	136	Gene No.
HFCEB37	НЕ8ЕҮ43	HE2GT20	HCWHZ24	HCQAI40	HFKFJ07	HBXGK12	.cDNA Clone ID
209008 04/28/97 209084	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209010 04/28/97 209085 05/29/97	209008 04/28/97 209084 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	ZAP Express	Lambda ZAP II	Uni-ZAP XR	ZAP Express	Vector
152	151	150	149	148	147	146	XEO NO: NO:
802	2399	2890	1405	734	1183	4313	Total NT Seq.
352	1181	150 2890 1178		Ļ.	-	1153	5' NT of Clone Seq.
802	2399		1405	734	1183	1153 4313	3' NT of Clone Seq.
	1265		801	285	149	1313	of Start
487	1265	1178	108	285	149	1313	of AA First First SEQ AA AA of ID of Signal NO: Sig Pep Y Pep
375	374	373	372	371	370	369	Y.O.SEQ SEQ SEQ
,	-	-	_	-	_	_	First AA of Sig Pep
	30	32	34		4	- 5	Las AA Of Sig
	31	32	35		42	19	First AA Last of AA Secreted of Portion ORF
<u></u>	34	39	63	19	254	42	Last AA ORF

149	148	147	146	145	144	143		Gene No.
HLMMU76	HKLAB16	HUSIT49	нјаа изб	HHGBR15	HGLAM46	HFTCT67		cDNA Clone ID
209008 04/28/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	05/29/97	ATCC Deposit Nr and Date
Lambda ZAP II	Lambda ZAP II	pSportl	pBluescript SK-	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR		Vector
159	158	157	156	155	154	153		× N B S N
1687	1625	2127	1251	642	154 2388	461		Total NT Seq.
1307	817	247	583	322	818	24		5' NT of Clone Seq.
1687	1625	2127	1251	642	2388	461		5' NT 3' NT of Clone Clone Seq. Seq.
1296	1012	383		400	648	145		5' NT of Start Codor
1296	1012	383	933	400	648	145		of First AA of Signal Pep
382	381	380	379	378	377	376		≺ <u>ö</u> ₽88
_	L	-	-	_		-		First AA of Sig Pep
28	18	47	16			37		Last AA of Sig Pep
29	19	48	17			<u>အ</u>		First AA of Secreted Portion
28	20	83	16	4		63		Last AA of ORF

										<u></u>	
157	156	156	155	154	153	152	151	150		Gene No.	
H6EAE26	HSKCP69	HSKCP69	HPTRC15	HOECU83	HNHFQ63	HNHEJ88	HNHED86	HMSKQ35		cDNA Clone ID	
209009	209009 04/28/97	209009 04/28/97	209009 04/28/97	209009 04/28/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209084 05/29/97	Nr and Date	ATCC
209009 Uni-ZAP XR 167	Uni-ZAP XR	Uni-ZAP XR	pBluescript	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
167	230	166	165	164	163	162	161	160		׊E	SEQ
882	1250	1251	2153	164 1400	753	519	770	1842		NT Seq.	
48	223		594	189	1			172		Seq.	of IN S
882	1250	1120	2153	1400	753	519	770	1463		Seq. Seq.	5' NT 3' NT of
155	393				164	242	30	319		Start Codon	C)
155			611	508	164	242	30	319		Signal NO: Pep Y	5' NT of First
390	453	389	388	387	386	385	384	383		≺ÖĘ	AA SEQ
	1	-	-	-	-	1	1	1		or Sig Pep	First AA
33	32			22	17	17	31	30		Sig Pep	Last AA
34	33			23	18	18	32	31		or Secreted Portion	First AA
153	171		13	33	67	24	46	33		윉유	

												0			_
168	167	166	165	164	163	162	161	160	159	158		Gene No.			_
HCFNF11	HCEZS40	HCEQA68	HCDDB78	HBMVP04	НВМТҮ28	HBHAD12	HAUAE83	HAICP19	HAGDQ47	HAGBX03		cDNA Clone ID			
209010	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209009 04/28/97	04/28/97	Nr and Date	Deposit	ATCC								
pSport1	Uni-ZAP XR	Uni-ZAP XR 176 1348	Uni-ZAP XR		Vector										
178	177	176	175	174	173	172	171	170	169	168		×Ö	Ħ,	SEO	i
1637	1502	1348	2379	888	1758	786	2003	1624	1307	1208		Seq.	Total		·
26	178		750	330	962	1	889	89	ı	1		Seq.	Clone	of N	
1607	1502	1348	2379	862	1758	786	2003	1483	1307	1208		Seq.	Clone Clone	of of of	
152	315	12	901		1184		1080	128	44	182		Start Codor	of	5' NT	
152	315	12	901	546	1184	176	1080	128	44	182		Signal Pep	AA of	First	5' NT
401	400	399	398	397	396	395	394	393	392	391		۲Ö	U,	SEO	
-	1	-	-	1		1	1		1	1		Sig Pep		AA	
44		28	18		27	17		18	22			Sig Pep			_
45		29	19		28	18		19	23			Secreted Portion	of,	First AA	
257	20	78	24	2	34	23	23	446	60	80		ORF of	₹	Last	

173	172	171	170	169	169		Gene No.
HE8MG65	HE2CT29	HDSAP81	HCUBL62	HCRBL20	HCRBL20		cDNA Clone ID
209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	04/28/97 209085 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	ZAP Express	Uni-ZAP XR	Uni-ZAP XR		Vector
183	182	181	180	231	179		× Ö B SEÖ N
2276	1128	968	519	1811	2911		Total NT Seq.
48	-	320	-	20	1103		5' NT of Clone Seq.
2276	1128	968	519	1811	2858		5' NT 3' NT of of Clone Clone Seq. Seq.
88	111	476	57	93	192		5' NT of Start Codor
88	111	476	57	93	192		of AA I of SEQ AA of ID Signal NO:
406	405	404	403	454	402		YO. BA
-	-	_	-	-	_		First AA of Sig Pep
37	26	27	28	36	32		Last AA of Sig Pep
38	27	28	29	37	33		First AA of Secreted Portion
257	94	79	32	95	424		Last AA of ORF

178	177	176	175	175	174	173	Gene No.
HETAR54	HEMDX17	HEMCV19	HEMAM41	HEMAM41	HE9FB42	HE8MG65	cDNA Clone ID
209010 04/28/97 209085	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Vector
188	187	186	233	185			× N D SEN
1848	654	941	1338	1337	2500	2271	Total NT Seg.
454		33	· · · · · · · · · · · · · · · · · · ·	8			5' NT of Clone Seq.
1848	654	931	1327	1328	1693	2232	5' NT 3' NT of of Total Clone Clone NT Seq. Seq.
948	137	79	175	175	810	79	5' NT of Start Codon
948	13/	79	175	C/1	910		of First AAA of Signal Pep
411	410	409	456	408	5	\$ 5	YNDSS SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ S
	-		-	-		-	First AA of Sig Pep
4		2.5	32	39	-	<u>.</u> .	Last AA of Sig Pep
15	10	24	33	<u> </u>	,	‡	First AA of Secreted Portion
202	33	1/8	170	140		623	Last AA of ORF

187	186	185	184	183	182	181	180	179		Gene No.
HHPSD37	HHPDW05	HHLBA89	HGLAM56	HGBF079	HFXHN68	HFKF140	HFGAB48	HETBX14		. cDNA Clone ID
209011 04/28/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	05/29/97	ATCC Deposit Nr and Date
pBluescript	Uni-ZAP XR	pBluescript SK-	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector
197	196	195	194	193	192	191	190	189		× Ö B Ö N
1282	1443	1001	1098	1538	2118	1941	906	1146		Total NT Seq.
66	-	1	68		777	120	,	157		5' NT of Clone Seq.
1282	1443	1001	1098	1538	2118	1002	906	1146		5' NT 3' NT of of Clone Clone Seq. Seq.
171	246	324		273	966	213				S' NT of Start Codor
171	246	324	185	273	966	213	245	74		of of First AA of Signal Pep
420	419	418	417	416	415	414	413	412		ΥÖ. BÖ SEÖ SEÖ
	-	1	1	-	1					First AA of Sig Pep
19	.21	25	28	23	23	18	30	14		Last AA of Sig Pep
20	22	26	29	24	24	19	31	15		First AA of Secreted Portion
37	21	39	69	49	50	218	32	53		Last AA of ORF

200	199	198	197	196	195	194	193	192	191	190	189	188	Gene No.		·
HNFAH08	HMSHQ24	HMSHM43	HLTDB65	HLTCY93	HLMIW92	HLHTC70	HLHSK94	нјрвв39	HJABZ65	HIASB53	HHSAK25	HHPSF70	cDNA Clone ID		
209011 04/28/97	Nr and Date	Deposit	ATCC												
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR 207	Uni-ZAP XR	Lambda ZAP II	pBluescript	pBluescript	Uni-ZAP XR	pBluescript SK-	pBluescript	Uni-ZAP XR	pBluescript	Vector		
210	209	208	207	206	205	204	203	202	201	200	199	198	×ö	Ħź	GN,
2110	1779	872	1480	2465	721	1057	203 1974	1617	779	200 1707	1740	951	NT Seq.	Total	
592	16	_	-	988	-	229	-	188	_	401	1390	26	Seq.	Clone	5' NT
2110	1779	872	1480	2465	721	1057	1794	1605	779	1195	1740			Clone Clone	3' NT
611	148	35		1225	244	365	112	182	23	652	1534		Start Codon	of	ላ: 2 ገ
61	148	35	371	1225	244	365	112	182	23	652	1534	162	Signal Pep	AA of	First 2
433	432	431	430	429	428	427	426	425	424	423	422	421	ΥĊ		£8
_	_		_	_	-	_	_	_	-	-	_	_	Sig Pep		First
18	24	~	15		25	23	26	28	26	26	19	16			Last
19	25	19	16		26	24	27	29	27	27	20	17	Secreted Portion	of	Firet AA
191	36	36	143	4	46	22	379	91	68	126	31	34			1 24

						T			
207	206	205	204	203	202	201	Gene No.		
HCDE095	нрнаС88	HOSFM22	НИНСМ59	91ZVHNH	HNGBE45	HNGAO10	cDNA Clone ID	•	
209007 04/28/97 209083 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	17	209011 04/28/97	209011 04/28/97	209011 04/28/97	Nr and Date	Deposit	
Uni-ZAP XR 217	Uni-ZAP XR 216 1705 384	Uni-ZAP XK	Uni-ZAP XR	Uni-ZAP XR 213 99/	Uni-ZAP XR 212 1551	Uni-ZAP XR	Vector		-
217	216	C17	214	213	212	211	×ö		
999	1705	1308	1496	/66	1551	938	Seq.	Total	
608	384	001				-	Seq.	Clone	5' NT
999			1300	1 49/	1001	938	Seq.	Clone	3' NT
273				707		$\overline{}$	NT Seq. Seq. Start Signal NO: Sig Sig Seq. Codon Pep Y Pep Pep	of :	۲۵ ا
273	549	600	200	707	1 1		Signal Pep	AA of	of First
440	434		478	127	436	404	¥.C	B	SEO A
-	-		1	-	-		Sig Pep	်င္	First
2	3 8) i	5	36	24) /	Pep	. č	A E
25	3	24	į	29	25	2) 6	Portion	of	First AA
ý	2 2	24		4	36	3 8	ORF S	\$ }	Last

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Table 1 summarizes the information corresponding to each "Gene No." described above. The nucleotide sequence identified as "NT SEQ ID NO:X" was assembled from partially homologous ("overlapping") sequences obtained from the "cDNA clone ID" identified in Table 1 and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous sequence of high redundancy (usually three to five overlapping sequences at each nucleotide position), resulting in a final sequence identified as SEQ ID NO:X.

The cDNA Clone ID was deposited on the date and given the corresponding deposit number listed in "ATCC Deposit No:Z and Date." Some of the deposits contain multiple different clones corresponding to the same gene. "Vector" refers to the type of vector contained in the cDNA Clone ID.

"Total NT Seq." refers to the total number of nucleotides in the contig identified by "Gene No." The deposited clone may contain all or most of these sequences, reflected by the nucleotide position indicated as "5' NT of Clone Seq." and the "3' NT of Clone Seq." of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the putative start codon (methionine) is identified as "5' NT of Start Codon." Similarly, the nucleotide position of SEQ ID NO:X of the predicted signal sequence is identified as "5' NT of First AA of Signal Pep."

The translated amino acid sequence, beginning with the methionine, is identified as "AA SEQ ID NO:Y," although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by these alternative open reading frames are specifically contemplated by the present invention.

The first and last amino acid position of SEQ ID NO:Y of the predicted signal peptide is identified as "First AA of Sig Pep" and "Last AA of Sig Pep." The predicted first amino acid position of SEQ ID NO:Y of the secreted portion is identified as "Predicted First AA of Secreted Portion." Finally, the amino acid position of SEQ ID NO:Y of the last amino acid in the open reading frame is identified as "Last AA of ORF."

SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in the deposited clone. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified in Table 1.

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Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

Accordingly, for those applications requiring precision in the nucleotide sequence or the amino acid sequence, the present invention provides not only the generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA containing a human cDNA of the invention deposited with the ATCC, as set forth in Table 1. The nucleotide sequence of each deposited clone can readily be determined by sequencing the deposited clone in accordance with known methods. The predicted amino acid sequence can then be verified from such deposits. Moreover, the amino acid sequence of the protein encoded by a particular clone can also be directly determined by peptide sequencing or by expressing the protein in a suitable host cell containing the deposited human cDNA, collecting the protein, and determining its sequence.

The present invention also relates to the genes corresponding to SEQ ID NO:X, SEQ ID NO:Y, or the deposited clone. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

Also provided in the present invention are species homologs. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for the desired homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below).

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It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the invention also can be purified from natural or recombinant sources using antibodies of the invention raised against the secreted protein in methods which are well known in the art.

Signal Sequences

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Methods for predicting whether a protein has a signal sequence, as well as the cleavage point for that sequence, are available. For instance, the method of McGeoch, Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 indicates the amino terminus of the secreted protein. The accuracy of predicting the cleavage points of known mammalian secretory proteins for each of these methods is in the range of 75-80%. (von Heinje, supra.) However, the two methods do not always produce the same predicted cleavage point(s) for a given protein.

In the present case, the deduced amino acid sequence of the secreted polypeptide was analyzed by a computer program called SignalP (Henrik Nielsen et al., Protein Engineering 10:1-6 (1997)), which predicts the cellular location of a protein based on the amino acid sequence. As part of this computational prediction of localization, the methods of McGeoch and von Heinje are incorporated. The analysis of the amino acid sequences of the secreted proteins described herein by this program provided the results shown in Table 1.

As one of ordinary skill would appreciate, however, cleavage sites sometimes vary from organism to organism and cannot be predicted with absolute certainty. Accordingly, the present invention provides secreted polypeptides having a sequence shown in SEQ ID NO:Y which have an N-terminus beginning within 5 residues (i.e., + or - 5 residues) of the predicted cleavage point. Similarly, it is also recognized that in some cases, cleavage of the signal sequence from a secreted protein is not entirely

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uniform, resulting in more than one secreted species. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Moreover, the signal sequence identified by the above analysis may not necessarily predict the naturally occurring signal sequence. For example, the naturally occurring signal sequence may be further upstream from the predicted signal sequence. However, it is likely that the predicted signal sequence will be capable of directing the secreted protein to the ER. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

10 Polynucleotide and Polypeptide Variants

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"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. The query sequence may be an entire sequence shown in Table 1, the ORF (open reading frame), or any fragement specified as described herein.

As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the presence invention can be determined conventionally using known computer programs. A preferred method for determing the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. (1990) 6:237-245). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are:

Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization

Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty 0.05, Window Size=500 or the length of the subject nucleotide sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the results. This is becuase the FASTDB program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only bases outside the 5' and 3' bases of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

For example, a 90 base subject sequence is aligned to a 100 base query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignement of the first 10 bases at 5' end. The 10 unpaired bases represent 10% of the sequence (number of bases at the 5' and 3' ends not matched/total number of bases in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 bases were perfectly matched the final percent identity would be 90%. In another example, a 90 base subject sequence is compared with a 100 base query sequence. This time the deletions are internal deletions so that there are no bases on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only bases 5' and 3' of the subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a query

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amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

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As a practical matter, whether any particular polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequences shown in Table 1 or to the amino acid sequence encoded by deposited DNA clone can be determined conventionally using known computer programs. A preferred method for determing the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. (1990) 6:237-245). In a sequence alignment the query and subject sequences are either both nucleotide sequences or both amino acid sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB amino acid alignment are: Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence due to N- or Cterminal deletions, not because of internal deletions, a manual correction must be made to the results. This is becuase the FASTDB program does not account for N- and Cterminal truncations of the subject sequence when calculating global percent identity. For subject sequences truncated at the N- and C-termini, relative to the the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, only query residue positions outside the farthest N- and C-terminal residues of the subject sequence.

For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the Nterminus of the subject sequence and therefore, the FASTDB alignment does not show a matching/alignment of the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal deletions so there are no residues at the N- or Ctermini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequnce are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

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The variants may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as E. coli).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 (1993), reported variant KGF proteins having heparin binding activity even after

deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)

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Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over 3,500 individual IL-1a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." (See, Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

Thus, the invention further includes polypeptide variants which show substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

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The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

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Polynucleotide and Polypeptide Fragments

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In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in the deposited clone or shown in SEQ ID NO:X. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in the deposited clone or the nucleotide sequence shown in SEQ ID NO:X. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, 500, 600, 2000 nucleotides) are preferred.

Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, 701-750, 751-800, 800-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, or 2001 to the end of SEQ ID NO:X or the cDNA contained in the deposited clone. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity. More preferably, these polynucleotides can be used as probes or primers as discussed herein.

In the present invention, a "polypeptide fragment" refers to a short amino acid sequence contained in SEQ ID NO:Y or encoded by the cDNA contained in the deposited clone. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, or 161 to the end of the coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the

carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

Particularly, N-terminal deletions of the polypeptide of the present invention can be described by the general formula m-p, where p is the total number of amino acids in the polypeptide and m is an integer from 2 to (p-1), and where both of these integers (m & p) correspond to the position of the amino acid residue identified in SEQ ID NO:Y.

Moreover, C-terminal deletions of the polypeptide of the present invention can also be described by the general formula 1-n, where n is an integer from 2 to (p-1), and again where these integers (n & p) correspond to the position of the amino acid residue identified in SEQ ID NO:Y.

The invention also provides polypeptides having one or more amino acids deleted from both the amino and the carboxyl termini, which may be described generally as having residues m-n of SEQ ID NO:Y, where m and n are integers as described above.

Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions.

Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

Epitopes & Antibodies

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In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an

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epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998-4002 (1983).)

Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')2 fragments) which are capable of specifically binding to protein. Fab and F(ab')2 fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred, as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

Fusion Proteins

Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the

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polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

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Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. Biochem. 270:3958-3964 (1995).)

Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. (EP-A 0232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D.

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Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. Chem. 270:9459-9471 (1995).)

Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a peptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).)

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

15 Vectors, Host Cells, and Protein Production

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

The polynucleotide insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance

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genes for culturing in E. coli and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as E. coli, Streptomyces and Salmonella typhimurium cells; fungal cells, such as yeast cells; insect cells such as Drosophila S2 and Spodoptera Sf9 cells; animal cells such as CHO, COS, 293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan.

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Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., Basic Methods In Molecular Biology (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

A polypeptide of this invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification.

Polypeptides of the present invention, and preferably the secreted form, can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein

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after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

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Uses of the Polynucleotides

Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes known techniques.

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The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each polynucleotide of the present invention can be used as a chromosome marker.

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Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment.

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Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted chromosomes, and preselection by hybridization to construct chromosome specific-cDNA libraries.

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Precise chromosomal location of the polynucleotides can also be achieved using fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

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For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes). Preferred polynucleotides correspond to the noncoding regions of the cDNAs because the coding sequences are

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more likely conserved within gene families, thus increasing the chance of cross hybridization during chromosomal mapping.

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Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library).) Assuming 1 megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker.

In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Both methods rely on binding of the polynucleotide to DNA or RNA. For these techniques, preferred polynucleotides are usually 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model

systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease.

Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell.

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The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for RFLP.

The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of

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unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers specific to particular tissue prepared from the sequences of the present invention. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination.

In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

Uses of the Polypeptides

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Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell . Biol. 105:3087-3096 (1987).) Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (125I, 121I), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technetium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

In addition to assaying secreted protein levels in a biological sample, proteins can also be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 131I, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic

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resonance, is introduced (for example, parenterally, subcutaneously, or intraperitoneally) into the mammal. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982).)

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Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an individual; (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

Moreover, polypeptides of the present invention can be used to treat disease. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B), to inhibit the activity of a polypeptide (e.g., an oncogene), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth).

Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

At the very least, the polypeptides of the present invention can be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

Biological Activities

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The polynucleotides and polypeptides of the present invention can be used in assays to test for one or more biological activities. If these polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides could be used to treat the associated disease.

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Immune Activity

A polypeptide or polynucleotide of the present invention may be useful in treating deficiencies or disorders of the immune system, by activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune deficiencies or disorders may be genetic, somatic, such as cancer or some autoimmune disorders, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, a polynucleotide or polypeptide of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

A polynucleotide or polypeptide of the present invention may be useful in treating or detecting deficiencies or disorders of hematopoietic cells. A polypeptide or polynucleotide of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat those disorders associated with a decrease in certain (or many) types hematopoietic cells. Examples of immunologic deficiency syndromes include, but are not limited to: blood protein disorders (e.g. agammaglobulinemia, dysgammaglobulinemia), ataxia telangiectasia, common variable immunodeficiency, Digeorge Syndrome, HIV infection, HTLV-BLV infection, leukocyte adhesion deficiency syndrome, lymphopenia, phagocyte bactericidal dysfunction, severe combined immunodeficiency (SCIDs), Wiskott-Aldrich Disorder, anemia, thrombocytopenia, or hemoglobinuria.

Moreover, a polypeptide or polynucleotide of the present invention could also be used to modulate hemostatic (the stopping of bleeding) or thrombolytic activity (clot formation). For example, by increasing hemostatic or thrombolytic activity, a polynucleotide or polypeptide of the present invention could be used to treat blood coagulation disorders (e.g., afibrinogenemia, factor deficiencies), blood platelet disorders (e.g. thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, a polynucleotide or polypeptide of the present invention that can

decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment of heart attacks (infarction), strokes, or scarring.

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A polynucleotide or polypeptide of the present invention may also be useful in treating or detecting autoimmune disorders. Many autoimmune disorders result from inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing autoimmune disorders.

Examples of autoimmune disorders that can be treated or detected by the present invention include, but are not limited to: Addison's Disease, hemolytic anemia, antiphospholipid syndrome, rheumatoid arthritis, dermatitis, allergic encephalomyelitis, glomerulonephritis, Goodpasture's Syndrome, Graves' Disease, Multiple Sclerosis, Myasthenia Gravis, Neuritis, Ophthalmia, Bullous Pemphigoid, Pemphigus, Polyendocrinopathies, Purpura, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Thyroiditis, Systemic Lupus Erythematosus, Autoimmune Pulmonary Inflammation, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye disease.

Similarly, allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated by a polypeptide or polynucleotide of the present invention. Moreover, these molecules can be used to treat anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

A polynucleotide or polypeptide of the present invention may also be used to treat and/or prevent organ rejection or graft-versus-host disease (GVHD). Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. The administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD.

Similarly, a polypeptide or polynucleotide of the present invention may also be used to modulate inflammation. For example, the polypeptide or polynucleotide may inhibit the proliferation and differentiation of cells involved in an inflammatory response. These molecules can be used to treat inflammatory conditions, both chronic and acute conditions, including inflammation associated with infection (e.g., septic

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shock, sepsis, or systemic inflammatory response syndrome (SIRS)), ischemiareperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, or resulting from over production of cytokines (e.g., TNF or IL-1.)

Hyperproliferative Disorders

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A polypeptide or polynucleotide can be used to treat or detect hyperproliferative disorders, including neoplasms. A polypeptide or polynucleotide of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, a polypeptide or polynucleotide of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

Examples of hyperproliferative disorders that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but are not limited to neoplasms located in the: abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

Similarly, other hyperproliferative disorders can also be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstron's Macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

Infectious Disease

A polypeptide or polynucleotide of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases

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may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, the polypeptide or polynucleotide of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

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Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus, Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza), Papovaviridae, Parvoviridae. Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g., Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiollitis, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), meningitis, opportunistic infections (e.g., AIDS). pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.

Similarly, bacterial or fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacterial families and fungi: Actinomycetales (e.g., Corynebacterium, Mycobacterium, Norcardia), Aspergillosis, Bacillaceae (e.g., Anthrax, Clostridium), Bacteroidaceae, Blastomycosis, Bordetella, Borrelia, Brucellosis, Candidiasis, Campylobacter, Coccidioidomycosis, Cryptococcosis, Dermatocycoses, Enterobacteriaceae (Klebsiella, Salmonella, Serratia, Yersinia), Erysipelothrix, Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales, Neisseriaceae (e.g., Acinetobacter, Gonorrhea, Menigococcal), Pasteurellacea Infections (e.g., Actinobacillus, Heamophilus, Pasteurella), Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, and Staphylococcal. These bacterial or fungal families can cause the following diseases or symptoms, including, but not limited to: bacteremia, endocarditis, eye infections (conjunctivitis, tuberculosis, uveitis), gingivitis, opportunistic infections (e.g., AIDS

any of these symptoms or diseases.

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related infections), paronychia, prosthesis-related infections, Reiter's Disease, respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, Dysentery, Paratyphoid Fever, food poisoning, Typhoid, pneumonia, Gonorrhea, meningitis, Chlamydia, Syphilis, Diphtheria, Leprosy, Paratuberculosis, Tuberculosis, Lupus, Botulism, gangrene, tetanus, impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections. A polypeptide or polynucleotide of the present invention can be used to treat or detect

Moreover, parasitic agents causing disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following families: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, Leishmaniasis, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas.

These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), Malaria, pregnancy complications, and toxoplasmosis. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.

Preferably, treatment using a polypeptide or polynucleotide of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

Regeneration

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A polynucleotide or polypeptide of the present invention can be used to differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteocarthritis, periodontal disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

Tissues that could be regenerated using the present invention include organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal

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or cardiac), vascular (including vascular endothelium), nervous, hematopoietic, and skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

Moreover, a polynucleotide or polypeptide of the present invention may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. A polynucleotide or polypeptide of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

Similarly, nerve and brain tissue could also be regenerated by using a polynucleotide or polypeptide of the present invention to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stoke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotide or polypeptide of the present invention.

Chemotaxis

A polynucleotide or polypeptide of the present invention may have chemotaxis activity. A chemotaxic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

A polynucleotide or polypeptide of the present invention may increase chemotaxic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotaxic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

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It is also contemplated that a polynucleotide or polypeptide of the present invention may inhibit chemotactic activity. These molecules could also be used to treat disorders. Thus, a polynucleotide or polypeptide of the present invention could be used as an inhibitor of chemotaxis.

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Binding Activity

A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from mammals, yeast, Drosophila, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule.

The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a

standard.

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Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

Other Activities

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A polypeptide or polynucleotide of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

A polypeptide or polynucleotide of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide or polynucleotide of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

A polypeptide or polynucleotide of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, caricadic rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity), hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

A polypeptide or polynucleotide of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

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Other Preferred Embodiments

Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Clone Sequence and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Start Codon and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Similarly preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ ID NO:X beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X.

Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.

Also preferred is a composition of matter comprising a DNA molecule which comprises a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1 for said cDNA Clone Identifier.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in the nucleotide sequence of a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the ATCC Deposit Number shown in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said human cDNA clone.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is a method for detecting in a biological sample a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method

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comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95%

identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

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Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1.

Also preferred is a polypeptide, wherein said sequence of contiguous amino acids is included in the amino acid sequence of SEQ ID NO:Y in the range of positions beginning with the residue at about the position of the First Amino Acid of the Secreted Portion and ending with the residue at about the Last Amino Acid of the Open Reading Frame as set forth for SEQ ID NO:Y in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a secreted portion of the secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is an isolated antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method comprises a step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group and determining whether the sequence of said polypeptide molecule in said sample is at least 90% identical to said sequence of at least 10 contiguous amino acids.

Also preferred is the above method wherein said step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of polypeptides in said sample to an antibody which binds specifically to a polypeptide

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comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

Also preferred is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is the above method for identifying the species, tissue or cell type of a biological sample, which method comprises a step of detecting polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

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Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said nucleotide sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

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Also preferred is an isolated nucleic acid molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a secreted portion of a human secreted protein comprising an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y beginning with the residue at the position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y wherein Y is an integer set forth in Table 1 and said position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y is defined in Table 1; and an amino acid sequence of a secreted portion of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The isolated polypeptide produced by this method is also preferred.

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Also preferred is a method of treatment of an individual in need of an increased level of a secreted protein activity, which method comprises administering to such an individual a pharmaceutical composition comprising an amount of an isolated polypeptide, polynucleotide, or antibody of the claimed invention effective to increase the level of said protein activity in said individual.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

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Examples

Example 1: Isolation of a Selected cDNA Clone From the Deposited Sample

Each cDNA clone in a cited ATCC deposit is contained in a plasmid vector. Table 1 identifies the vectors used to construct the cDNA library from which each clone was isolated. In many cases, the vector used to construct the library is a phage vector from which a plasmid has been excised. The table immediately below correlates the related plasmid for each phage vector used in constructing the cDNA library. For example, where a particular clone is identified in Table 1 as being isolated in the vector "Lambda Zap," the corresponding deposited clone is in "pBluescript."

	Vector Used to Construct Library	Corresponding Deposited Plasmid
	Lambda Zap	pBluescript (pBS)
	Uni-Zap XR	pBluescript (pBS)
	Zap Express	pBK
25	lafmid BA	plafmid BA
	pSport1	pSport1
	pCMVSport 2.0	pCMVSport 2.0
	pCMVSport 3.0	pCMVSport 3.0
	pCR [®] 2.1	pCR [®] 2.1

Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res. 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., Nucleic Acids Res. 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1

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Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS. The S and K refers to the orientation of the polylinker to the T7 and T3 primer sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which are the first sites on each respective end of the linker). "+" or "-" refer to the orientation of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the f1 ori generates sense strand DNA and in the other, antisense.

Vectors pSport1, pCMVSport 2.0 and pCMVSport 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1 Blue. Vector pCR®2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991).) Preferably, a polynucleotide of the present invention does not comprise the phage vector sequences identified for the particular clone in Table 1, as well as the corresponding plasmid vector sequences designated above.

The deposited material in the sample assigned the ATCC Deposit Number cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 50 plasmid DNAs, each containing a different cDNA clone; but such a deposit sample may include plasmids for more or less than 50 cDNA clones, up to about 500 cDNA clones.

Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO:X.

Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ³²P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).)

The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993).)

Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase should then be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

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Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the cDNA sequence corresponding to SEQ ID NO:X., according to the method described in Example 1. (See also, Sambrook.)

Example 3: Tissue Distribution of Polypeptide

Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method described in Example 1 is labeled with P³² using the rediprimeTM DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN-100TM column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for mRNA expression.

Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) (Clontech) are examined with the labeled probe using ExpressHybTM hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are mounted and exposed to film at -70°C overnight, and the films developed according to standard procedures.

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Example 4: Chromosomal Mapping of the Polynucleotides

An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of conditions: 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions is analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

Example 5: Bacterial Expression of a Polypeptide

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A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Amp^r), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kan^r). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D.⁶⁰⁰) of between 0.4 and 0.6. IPTG

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(Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

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Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., *supra*).

Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM immidazole. Immidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

In addition to the above expression vector, the present invention further includes an expression vector comprising phage operator and promoter elements operatively linked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession Number 209645, deposited on February 25, 1998.) This vector contains: 1) a neomycinphosphotransferase gene as a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgarno sequence, and 6) the lactose operon repressor gene (lacIq). The origin of replication (oriC) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter sequence and operator sequences are made synthetically.

DNA can be inserted into the pHEa by restricting the vector with NdeI and XbaI, BamHI, XhoI, or Asp718, running the restricted product on a gel, and isolating the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA

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insert is generated according to the PCR protocol described in Example 1, using PCR primers having restriction sites for NdeI (5' primer) and XbaI, BamHI, XhoI, or Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

Example 6: Purification of a Polypeptide from an Inclusion Body

The following alternative method can be used to purify a polypeptide expressed in *E coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells are then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with $0.16\,\mu m$ membrane filter with appropriate surface area

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(e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded.

The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

Example 7: Cloning and Expression of a Polypeptide in a Baculovirus Expression System

In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak Drosophila promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

Many other baculovirus vectors can be used in place of the vector above, such as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-39 (1989).

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Specifically, the cDNA sequence contained in the deposited clone, including the AUG initiation codon and the naturally associated leader sequence identified in Table 1, is amplified using the PCR protocol described in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.).

The fragment and the dephosphorylated plasmid are ligated together with T4 DNA ligase. E. coli HB101 or other suitable E. coli hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.

Five μg of a plasmid containing the polynucleotide is co-transfected with 1.0 μg of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One μg of BaculoGold™ virus DNA and 5 μg of the plasmid are mixed in a sterile well of a microtiter plate containing 50 μl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm

tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

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After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, supra. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 µl of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 μCi of ³⁵Smethionine and 5 µCi 35S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

30 Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates

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the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSport 2.0, and pCMVSport 3.0. Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No.209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985).) Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the

polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the vector does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene is used for transfection. Five µg of the expression plasmid pC6 is cotransfected with 0.5 µg of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of metothrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μM , 2 μM , 5 μM , 10 m M, 20 m M). The same procedure is repeated until clones are obtained which grow at a concentration of 100 - $200 \, \mu M$. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

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Example 9: Protein Fusions

The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394,827; Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halflife time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.

For example, if pC4 (Accession No. 209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

If the naturally occurring signal sequence is used to produce the secreted protein, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

Human IgG Fc region:

GGGATCCGGAGCCCAAATCTTCTGACAAAACTCACACATGCCCACCGTGCC CAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCCCCCAAAACC CAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGTGGT GGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACG GCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAAC

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AGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGGCTG
AATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAACCCCC
ATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGT
GTACACCCTGCCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCT
GACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAGTGGGA
GAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGG
ACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCA
GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGC
ACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGAGTGC
GACGGCCGCGACTCTAGAGGAT (SEQ ID NO:1)

Example 10: Production of an Antibody from a Polypeptide

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The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) For example, cells expressing a polypeptide of the present invention is administered to an animal to induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of the secreted protein is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In the most preferred method, the antibodies of the present invention are monoclonal antibodies (or protein binding fragments thereof). Such monoclonal antibodies can be prepared using hybridoma technology. (Köhler et al., Nature 256:495 (1975); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J. Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures involve immunizing an animal (preferably a mouse) with polypeptide or, more preferably, with a secreted polypeptide-expressing cell. Such cells may be cultured in any suitable tissue culture medium; however, it is preferable to culture cells in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100 μg/ml of streptomycin.

The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as

described by Wands et al. (Gastroenterology 80:225-232 (1981).) The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide.

Alternatively, additional antibodies capable of binding to the polypeptide can be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and can be used to immunize an animal to induce formation of further protein-specific antibodies.

It will be appreciated that Fab and F(ab')2 and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, secreted protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For in vivo use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

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Example 11: Production Of Secreted Protein For High-Throughput Screening Assays

The following protocol produces a supernatant containing a polypeptide to be tested. This supernatant can then be used in the Screening Assays described in Examples 13-20.

First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a

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working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) and incubate at RT for 20 minutes. Be sure to distribute the solution over each well (note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The PBS should remain in the well until just prior to plating the cells and plates may be poly-lysine coated in advance for up to two weeks.

Plate 293T cells (do not carry cells past P+20) at 2 x 10⁵ cells/well in .5ml DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and L-glutamine (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

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The next day, mix together in a sterile solution basin: 300 ul Lipofectamine (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression vector containing a polynucleotide insert, produced by the methods described in Examples 8 or 9, into an appropriately labeled 96-well round bottom plate. With a multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well. Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a control, one plate of vector DNA lacking an insert should be transfected with each set of transfections.

Preferably, the transfection should be performed by tag-teaming the following tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too much time on PBS. First, person A aspirates off the media from four 24-well plates of cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off PBS rinse, and person B, using a12-channel pipetter with tips on every other channel, adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to the even wells, to each row on the 24-well plates. Incubate at 37°C for 6 hours.

While cells are incubating, prepare appropriate media, either 1%BSA in DMEM with 1x penstrep, or CHO-5 media (116.6 mg/L of CaCl2 (anhyd); 0.00130 mg/L CuSO₄-5H₂O; 0.050 mg/L of Fe(NO₃)₃-9H₂O; 0.417 mg/L of FeSO₄-7H₂O; 311.80 mg/L of Kcl; 28.64 mg/L of MgCl₂; 48.84 mg/L of MgSO₄; 6995.50 mg/L of NaCl; 2400.0 mg/L of NaHCO₃; 62.50 mg/L of NaH₂PO₄-H₂O; 71.02 mg/L of Na₂HPO4; .4320 mg/L of ZnSO₄-7H₂O; .002 mg/L of Arachidonic Acid; 1.022 mg/L of Cholesterol; .070 mg/L of DL-alpha-Tocopherol-Acetate; 0.0520 mg/L of Linoleic Acid; 0.010 mg/L of Linolenic Acid; 0.010 mg/L of Palmitric Acid; 0.010 mg/L of Palmitric Acid; 100 mg/L of

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Pluronic F-68; 0.010 mg/L of Stearic Acid; 2.20 mg/L of Tween 80; 4551 mg/L of D-Glucose; 130.85 mg/ml of L- Alanine; 147.50 mg/ml of L-Arginine-HCL; 7.50 mg/ml of L-Asparagine-H₂0; 6.65 mg/ml of L-Aspartic Acid; 29.56 mg/ml of L-Cystine-2HCL-H,0; 31.29 mg/ml of L-Cystine-2HCL; 7.35 mg/ml of L-Glutamic Acid; 365.0 mg/ml of L-Glutamine; 18.75 mg/ml of Glycine; 52.48 mg/ml of L-Histidine-HCL-H₂0; 106.97 mg/ml of L-Isoleucine; 111.45 mg/ml of L-Leucine; 163.75 mg/ml of L-Lysine HCL; 32.34 mg/ml of L-Methionine; 68.48 mg/ml of L-Phenylalainine; 40.0 mg/ml of L-Proline; 26.25 mg/ml of L-Serine; 101.05 mg/ml of L-Threonine; 19.22 mg/ml of L-Tryptophan; 91.79 mg/ml of L-Tryrosine-2Na-2H,0; 99.65 mg/ml of L-Valine; 0.0035 mg/L of Biotin; 3.24 mg/L of D-Ca Pantothenate; 11.78 mg/L of 10 Choline Chloride; 4.65 mg/L of Folic Acid; 15.60 mg/L of i-Inositol; 3.02 mg/L of Niacinamide; 3.00 mg/L of Pyridoxal HCL; 0.031 mg/L of Pyridoxine HCL; 0.319 mg/L of Riboflavin; 3.17 mg/L of Thiamine HCL; 0.365 mg/L of Thymidine; and 0.680 mg/L of Vitamin B₁₂; 25 mM of HEPES Buffer; 2.39 mg/L of Na Hypoxanthine; 15 0.105 mg/L of Lipoic Acid; 0.081 mg/L of Sodium Putrescine-2HCL; 55.0 mg/L of Sodium Pyruvate; 0.0067 mg/L of Sodium Selenite; 20uM of Ethanolamine; 0.122 mg/L of Ferric Citrate; 41.70 mg/L of Methyl-B-Cyclodextrin complexed with Linoleic Acid; 33.33 mg/L of Methyl-B-Cyclodextrin complexed with Oleic Acid; and 10 mg/L of Methyl-B-Cyclodextrin complexed with Retinal) with 2mm glutamine and 1x 20 penstrep. (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene conical.

The transfection reaction is terminated, preferably by tag-teaming, at the end of the incubation period. Person A aspirates off the transfection media, while person B adds 1.5ml appropriate media to each well. Incubate at 37°C for 45 or 72 hours depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours.

On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep well plate and the remaining supernatant into a 2ml deep well. The supernatants from each well can then be used in the assays described in Examples 13-20.

It is specifically understood that when activity is obtained in any of the assays described below using a supernatant, the activity originates from either the polypeptide directly (e.g., as a secreted protein) or by the polypeptide inducing expression of other proteins, which are then secreted into the supernatant. Thus, the invention further provides a method of identifying the protein in the supernatant characterized by an activity in a particular assay.

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Example 12: Construction of GAS Reporter Construct

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One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive responsive element ("ISRE"), located in the promoter of many genes. The binding of a protein to these elements alter the expression of the associated gene.

GAS and ISRE elements are recognized by a class of transcription factors called Signal Transducers and Activators of Transcription, or "STATs." There are six members of the STATs family. Stat1 and Stat3 are present in many cell types, as is Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in many cell types though it has been found in T helper class I, cells after treatment with IL-12. Stat5 was originally called mammary growth factor, but has been found at higher concentrations in other cells including myeloid cells. It can be activated in tissue culture cells by many cytokines.

The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks") family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarity and are generally catalytically inactive in resting cells.

The Jaks are activated by a wide range of receptors summarized in the Table below. (Adapted from review by Schidler and Darnell, Ann. Rev. Biochem. 64:621-51 (1995).) A cytokine receptor family, capable of activating Jaks, is divided into two groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and (b) Class 2 includes IFN-a, IFN-g, and IL-10. The Class 1 receptors share a conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a WSXWS motif (a membrane proxial region encoding Trp-Ser-Xxx-Trp-Ser (SEQ ID NO:2)).

Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn activate STATs, which then translocate and bind to GAS elements. This entire process is encompassed in the Jaks-STATs signal transduction pathway.

Therefore, activation of the Jaks-STATs pathway, reflected by the binding of the GAS or the ISRE element, can be used to indicate proteins involved in the proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway. (See Table below.) Thus, by using GAS elements linked to reporter molecules, activators of the Jaks-STATs pathway can be identified.

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	<u>Ligand</u>	tyk2	JAKs Jakl	Jak2	Jak3	<u>STATS</u>	GAS(elements) or ISRE
5	IFN family IFN-a/B IFN-g II-10	+	+ + ?	- + ?	- -	1,2,3 1 1,3	ISRE GAS (IRF1>Lys6>IFP)
10	gp130 family IL-6 (Pleiotrohic) II-1 (Pleiotrohic) OnM(Pleiotrohic)	+ ? ?	+ + + + .	+ ? +	? ? ?	1,3 1,3 1,3	GAS (IRF1>Lys6>IFP)
15	LIF(Pleiotrohic) CNTF(Pleiotrohic) G-CSF(Pleiotrohic) IL-12(Pleiotrohic)	? -/+ ? +	+ + + -	+ + ? +	? ? ? +	1,3 1,3 1,3 1,3	
20	g-C family IL-2 (lymphocytes) IL-4 (lymph/myeloid) IL-7 (lymphocytes) IL-9 (lymphocytes)	- - -	+ + +	-	+ + +	1,3,5 6 5 5	GAS GAS (IRF1 = IFP >>Ly6)(IgH) GAS GAS
25	IL-13 (lymphocyte) IL-15 gp140 family	?	+ +	?	?	6 5 .	GAS GAS
30	IL-3 (myeloid) IL-5 (myeloid) GM-CSF (myeloid)	- - -	- - -	++++++	- - -	5 5 5	GAS (IRF1>IFP>>Ly6) GAS GAS
25	Growth hormone fam GH PRL EPO	ily ? ? ?	- +/- -	+ + + +	-	5 1,3,5 5	GAS(B-CAS>IRF1=IFP>>Ly6)
35	Receptor Tyrosine Ki EGF PDGF CSF-1	nases ? ? ?	+ + +	+ + +	-	1,3 1,3	GAS (IRF1)
40	COI-1	;	7	T	-	1,3	GAS (not IRF1)

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To construct a synthetic GAS containing promoter element, which is used in the Biological Assays described in Examples 13-14, a PCR based strategy is employed to generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies of the GAS binding site found in the IRF1 promoter and previously demonstrated to bind STATs upon induction with a range of cytokines (Rothman et al., Immunity 1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5' primer also contains 18bp of sequence complementary to the SV40 early promoter sequence and is flanked with an XhoI site. The sequence of the 5' primer is: 5':GCGCCTCGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCG AAATGATTTCCCCGAAATATCTGCCATCTCAATTAG:3' (SEQ ID NO:3)

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The downstream primer is complementary to the SV40 promoter and is flanked with a Hind III site: 5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEO ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with Xhol/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing with forward and reverse primers confirms that the insert contains the following sequence:

5': CTCGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCGAAATG ATTTCCCCGAAATATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCC CTAACTCCGCCCATCCCGCCCTAACTCCGCCCAGTTCCGCCCATTCTCCGC CCCATGGCTGACTAATTTTTTTATTTATGCAGAGGCCGAGGCCGCCTCGGC CTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGGCTTT TGCAAAAAGCTT:3' (SEQ ID NO:5)

With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2 reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of SEAP, in this or in any of the other Examples. Well known reporter molecules that can be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any protein detectable by an antibody.

The above sequence confirmed synthetic GAS-SV40 promoter element is subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and XhoI, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter element, to create the GAS-SEAP vector. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

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Thus, in order to generate mammalian stable cell lines expressing the GAS-SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using SalI and NotI, and inserted into a backbone vector containing the neomycin resistance gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning site, to create the GAS-SEAP/Neo vector. Once this vector is transfected into mammalian cells, this vector can then be used as a reporter molecule for GAS binding as described in Examples 13-14.

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Other constructs can be made using the above description and replacing GAS with a different promoter sequence. For example, construction of reporter molecules containing NFK-B and EGR promoter sequences are described in Examples 15 and 16. However, many other promoters can be substituted using the protocols described in these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, Il-2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte.

Example 13: High-Throughput Screening Assay for T-cell Activity.

The following protocol is used to assess T-cell activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate T-cells. T-cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and Molt-4 cells (ATCC Accession No. CRL-1582) cells can also be used.

Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GAS-SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure described below). The transfected cells are seeded to a density of approximately 20,000 cells per well and transfectants resistant to 1 mg/ml genticin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of interferon gamma. The dose response of a selected clone is demonstrated.

Specifically, the following protocol will yield sufficient cells for 75 wells containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI + 10% serum with 1%Pen-Strep. Combine 2.5 mls of OPTI-MEM (Life Technologies)

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with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

During the incubation period, count cell concentration, spin down the required number of cells (10⁷ per transfection), and resuspend in OPTI-MEM to a final concentration of 10⁷ cells/ml. Then add 1ml of 1 x 10⁷ cells in OPTI-MEM to T25 flask and incubate at 37°C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum.

The Jurkat:GAS-SEAP stable reporter lines are maintained in RPMI + 10% serum, 1 mg/ml Genticin, and 1% Pen-Strep. These cells are treated with supernatants containing a polypeptide as produced by the protocol described in Example 11.

On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The exact number of cells required will depend on the number of supernatants being screened. For one 96 well plate, approximately 10 million cells (for 10 plates, 100 million cells) are required.

Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100, 000 cells per well).

After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12 channel pipette. In addition, a dose of exogenous interferon gamma (0.1, 1.0, 10 ng) is added to wells H9, H10, and H11 to serve as additional positive controls for the assay.

The 96 well dishes containing Jurkat cells treated with supernatants are placed in an incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples from each well are then transferred to an opaque 96 well plate using a 12 channel pipette. The opaque plates should be covered (using sellophene covers) and stored at -20°C until SEAP assays are performed according to Example 17. The plates containing the remaining treated cells are placed at 4°C and serve as a source of material for repeating the assay on a specific well if desired.

As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate Jurkat T cells. Over 30 fold induction is typically observed in the positive control wells.

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Example 14: High-Throughput Screening Assay Identifying Myeloid Activity

The following protocol is used to assess myeloid activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate myeloid cells. Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 12, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2x10e⁷ U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl, 375 uM Na₂HPO₄.7H₂O, 1 mM MgCl₂, and 675 uM CaCl₂. Incubate at 37°C for 45 min.

Wash the cells with RPMI 1640 medium containing 10% FBS and then resuspend in 10 ml complete medium and incubate at 37°C for 36 hr.

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 ug/ml G418 for couple of passages.

These cells are tested by harvesting $1x10^8$ cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of $5x10^5$ cells/ml. Plate 200 ul cells per well in the 96-well plate (or $1x10^5$ cells/well).

Add 50 ul of the supernatant prepared by the protocol described in Example 11. Incubate at 37°C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 17.

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Example 15: High-Throughput Screening Assay Identifying Neuronal Activity.

When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1 promoter linked to reporter molecules, activation of cells can be assessed.

Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat phenochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells can be assessed.

The EGR/SEAP reporter construct can be assembled by the following protocol. The EGR-1 promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871 (1991)) can be PCR amplified from human genomic DNA using the following primers:

5' GCGCTCGAGGGATGACAGCGATAGAACCCCGG -3' (SEQ ID NO:6) 5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO:7)

Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector using restriction enzymes XhoI/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1 promoter.

To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by growing the cells in 300 ug/ml G418. The G418-free medium is used for routine

growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

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The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as 5×10^5 cells/ml.

Add 200 ul of the cell suspension to each well of 96-well plate (equivalent to $1x10^5$ cells/well). Add 50 ul supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ul of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

Example 16: High-Throughput Screening Assay for T-cell Activity

NF-kB (Nuclear Factor kB) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF-kB regulates the expression of genes involved in immune cell activation, control of apoptosis (NF-kB appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

In non-stimulated conditions, NF- κB is retained in the cytoplasm with I- κB (Inhibitor κB). However, upon stimulation, I- κB is phosphorylated and degraded, causing NF- κB to shuttle to the nucleus, thereby activating transcription of target genes. Target genes activated by NF- κB include IL-2, IL-6, GM-CSF, ICAM-1 and class 1 MHC.

Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF-kB promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF-kB would be useful in treating

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diseases. For example, inhibitors of NF-kB could be used to treat those diseases related to the acute or chronic activation of NF-kB, such as rheumatoid arthritis.

To construct a vector containing the NF-κB promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF-κB binding site (GGGGACTTTCCC) (SEQ ID NO:8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site: 5':GCGGCCTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGGACTTTCCAACTTAG:3' (SEQ ID NO:9)

The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:

5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEO ID NO:4)

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PCR amplification is performed using the SV40 promoter template present in the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2-. (Stratagene)

Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

Next, replace the SV40 minimal promoter element present in the pSEAP2promoter plasmid (Clontech) with this NF-kB/SV40 fragment using XhoI and HindIII.
However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

In order to generate stable mammalian cell lines, the NF-κB/SV40/SEAP

cassette is removed from the above NF-κB/SEAP vector using restriction enzymes SalI and NotI, and inserted into a vector containing neomycin resistance. Particularly, the NF-κB/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with SalI and NotI.

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Once NF-kB/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 13. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 13. As a positive control, exogenous TNF alpha (0.1,1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

Example 17: Assay for SEAP Activity

As a reporter molecule for the assays described in Examples 13-16, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 μ l of 2.5x dilution buffer into Optiplates containing 35 μ l of a supernatant. Seal the plates with a plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven heating.

Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 µl Assay Buffer and incubate at room temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the table below). Add 50 µl Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each time and start the second set 10 minutes later.

Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

Meaction	butter rormulation:	
# of plates	Rxn buffer diluent (ml)	CSPD (ml)
10	60	3
11	65	3.25
12	70	3.5
13	75	3.75
14	80	4
15	85	4.25
16	90	4.5
17	95	4.75
18	100	5
19	105	5.25
20	110	5.5
21	115	5.75
22	120	6

23	125	6.25
24	130	6.5
25	135	6.75
26	140	7
27	145	7.25
28	150	7.5
29	155	7.75
30	160	8
31	165	8.25
32	170	8.5
33	175	8.75
34	180	9
35	185	9.25
36	190	9.5
37	195	9.75
38	200	10
39	205	10.25
40	210	10.5
41	215	10.75
42	220	11
43	225	11.25
44	230	11.5
45	235	11.75
46	240	12
47	245	12.25
48	250	12.5
49	255	12.75
50	260	13

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Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane potential. These alterations can be measured in an assay to identify supernatants which bind to receptors of a particular cell. Although the following protocol describes an assay for calcium, this protocol can easily be modified to detect changes in potassium, sodium, pH, membrane potential, or any other small molecule which is detectable by a fluorescent probe.

The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small molecules. Clearly, any fluorescent molecule detecting a small molecule can be used instead of the calcium fluorescent molecule, fluo-3, used here.

For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black 96-well plate with clear bottom. The plate is incubated in a CO₂ incubator for 20 hours. The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash.

A stock solution of 1 mg/ml fluo-3 is made in 10% pluronic acid DMSO. To load the cells with fluo-3, 50 ul of 12 ug/ml fluo-3 is added to each well. The plate is incubated at 37°C in a CO₂ incubator for 60 min. The plate is washed four times in the Biotek washer with HBSS leaving 100 ul of buffer.

For non-adherent cells, the cells are spun down from culture media. Cells are re-suspended to 2-5x10⁶ cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml fluo-3 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice with HBSS, resuspended to 1x10⁶ cells/ml, and dispensed into a microplate, 100 ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 ul, followed by an aspiration step to 100 ul final volume.

For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-3. The supernatant is added to the well, and a change in fluorescence is detected.

To measure the fluorescence of intracellular calcium, the FLIPR is set for the following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and (6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular signaling event which has resulted in an increase in the intracellular Ca⁺⁺ concentration.

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Example 19: High-Throughput Screening Assay Identifying Tyrosine Kinase Activity

The Protein Tyrosine Kinases (PTK) represent a diverse group of transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase RPTK) group are receptors for a range of mitogenic and metabolic growth factors including the PDGF, FGF, EGF, NGF, HGF and Insulin receptor subfamilies. In addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

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Activation of RPTK by ligands involves ligand-mediated receptor dimerization, resulting in transphosphorylation of the receptor subunits and activation of the cytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and non-receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors (e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

Because of the wide range of known factors capable of stimulating tyrosine kinase activity, the identification of novel human secreted proteins capable of activating tyrosine kinase signal transduction pathways are of interest. Therefore, the following protocol is designed to identify those novel human secreted proteins capable of activating the tyrosine kinase signal transduction pathways.

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Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or 10% Matrigel purchased from Becton Dickinson (Bedford,MA), or calf serum, rinsed with PBS and stored at 4°C. Cell growth on these plates is assayed by seeding 5,000 cells/well in growth medium and indirect quantitation of cell number through use of alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford,MA) are used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture plates can also be used in some proliferation experiments.

To prepare extracts, A431 cells are seeded onto the nylon membranes of Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium. Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na3VO4, 2 mM Na4P2O7 and a cocktail of protease inhibitors (# 1836170) obtained from Boeheringer Mannheim (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract filtered through the 0.45 mm membrane bottoms of each well using house vacuum. Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum manifold and immediately placed on ice. To obtain extracts clarified by centrifugation, the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

Test the filtered extracts for levels of tyrosine kinase activity. Although many methods of detecting tyrosine kinase activity are known, one method is described here.

Generally, the tyrosine kinase activity of a supernatant is evaluated by determining its ability to phosphorylate a tyrosine residue on a specific substrate (a

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biotinylated peptide). Biotinylated peptides that can be used for this purpose include PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for a range of tyrosine kinases and are available from Boehringer Mannheim.

The tyrosine kinase reaction is set up by adding the following components in order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg2+ (5mM ATP/50mM MgCl₂), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride, pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl₂, 5 mM MnCl₂, 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the components gently and preincubate the reaction mix at 30°C for 2 min. Initial the reaction by adding 10ul of the control enzyme or the filtered supernatant.

The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mm EDTA and place the reactions on ice.

Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This allows the streptavadin coated 96 well plate to associate with the biotinylated peptide. Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of anti-phospotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37°C for one hour. Wash the well as above.

Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and incubate at room temperature for at least 5 mins (up to 30 min). Measure the absorbance of the sample at 405 nm by using ELISA reader. The level of bound peroxidase activity is quantitated using an ELISA reader and reflects the level of tyrosine kinase activity.

Example 20: High-Throughput Screening Assay Identifying Phosphorylation Activity

As a potential alternative and/or compliment to the assay of protein tyrosine
kinase activity described in Example 19, an assay which detects activation
(phosphorylation) of major intracellular signal transduction intermediates can also be
used. For example, as described below one particular assay can detect tyrosine
phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other
molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase,

Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other

phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by substituting these molecules for Erk-1 or Erk-2 in the following assay.

Specifically, assay plates are made by coating the wells of a 96-well ELISA plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp, (RT). The plates are then rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this step can easily be modified by substituting a monoclonal antibody detecting any of the above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C until use.

A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and cultured overnight in growth medium. The cells are then starved for 48 hr in basal medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts filtered directly into the assay plate.

After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation.

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Example 21: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is be isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer solutions described in Sidransky, D., et al., Science 252:706 (1991).

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PCR products are then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies).

The intron-exon borders of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations is then cloned and sequenced to validate the results of the direct sequencing.

PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals are identified by mutations not present in unaffected individuals.

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Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'triphosphate (Boehringer Manheim), and FISH performed as described in Johnson, Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

15 Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. 20 et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated disease.

Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a **Biological Sample**

A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

For example, antibody-sandwich ELISAs are used to detect polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10.

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The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbounded polypeptide.

Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbounded conjugate.

Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

Example 23: Formulating a Polypeptide

The secreted polypeptide composition will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the secreted polypeptide alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of secreted polypeptide administered parenterally per dose will be in the range of about 1 μ g/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the secreted polypeptide is typically administered at a dose rate of about 1 μ g/kg/hour to about 50 μ g/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

Pharmaceutical compositions containing the secreted protein of the invention are administered orally, rectally, parenterally, intracistemally, intravaginally,

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intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

The secreted polypeptide is also suitably administered by sustained-release systems. Suitable examples of sustained-release compositions include semi-permeable polymer matrices in the form of shaped articles, e.g., films, or mirocapsules. Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (R. Langer et al., J. Biomed. Mater. Res. 15:167-277 (1981), and R. Langer, Chem. Tech. 12:98-105 (1982)), ethylene vinyl acetate (R. Langer et al.) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988). Sustained-release compositions also include liposomally entrapped polypeptides. Liposomes containing the secreted polypeptide are prepared by methods known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci. USA 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal secreted polypeptide therapy.

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For parenteral administration, in one embodiment, the secreted polypeptide is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to polypeptides.

Generally, the formulations are prepared by contacting the polypeptide uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

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The carrier suitably contains minor amounts of additives such as substances that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, manose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

The secreted polypeptide is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

Any polypeptide to be used for therapeutic administration can be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

Polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized polypeptide using bacteriostatic Water-for-Injection.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the present invention may be employed in conjunction with other therapeutic compounds.

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Example 24: Method of Treating Decreased Levels of the Polypeptide

It will be appreciated that conditions caused by a decrease in the standard or normal expression level of a secreted protein in an individual can be treated by administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a pharmaceutical composition comprising an amount of the polypeptide to increase the activity level of the polypeptide in such an individual.

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For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the polypeptide is in the secreted form. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 23.

Example 25: Method of Treating Increased Levels of the Polypeptide

Antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer.

For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, 2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is provided in Example 23.

Example 26: Method of Treatment Using Gene Therapy

One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin) is added. The flasks are then incubated at 37°C for approximately one week.

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At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is produced.

The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

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Example 27: Method of Treatment Using Gene Therapy - In Vivo

Another aspect of the present invention is using *in vivo* gene therapy methods to treat disorders, diseases and conditions. The gene therapy method relates to the introduction of naked nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into an animal to increase or decrease the expression of the polypeptide of the present invention. A polynucleotide of the present invention may be operatively linked to a promoter or any other genetic elements necessary for the expression of the encoded polypeptide by the target tissue. Such gene therapy and delivery techniques and methods are known in the art, see, for example, WO90/11092, WO98/11779; U.S. Patent NO. 5693622, 5705151, 5580859; Tabata H. et al. (1997) Cardiovasc. Res. 35(3):470-479, Chao J et al. (1997) Pharmacol. Res. 35(6):517-522, Wolff J.A. (1997) Neuromuscul. Disord. 7(5):314-318, Schwartz B. et al. (1996) Gene Ther. 3(5):405-411, Tsurumi Y. et al. (1996) Circulation 94(12):3281-3290 (incorporated herein by reference).

The polynucleotide constructs of the present invention may be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, intestine and the like). These polynucleotide constructs can be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

The term "naked" polynucleotide, DNA or RNA, refers to sequences that are free from any delivery vehicle that acts to assist, promote, or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotides may also be delivered in liposome formulations (such as those taught in Felgner P.L. et al. (1995) Ann. NY Acad. Sci. 772:126-139 and Abdallah B. et al. (1995) Biol. Cell 85(1):1-7) which can be prepared by methods well known to those skilled in the art.

The polynucleotide vector constructs of the present invention used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Any strong promoter known to those skilled in the art can be used for driving the expression of DNA. Unlike other gene therapies techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

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The polynucleotide construct of the present invention can be delivered to the interstitial space of tissues within the an animal, including of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. In vivo muscle cells are particularly competent in their ability to take up and express polynucleotides.

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For the naked polynucleotide injection, an effective dosage amount of DNA or RNA will be in the range of from about 0.05 g/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration. The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues, throat or mucous membranes of the nose. In addition, naked polynucleotide constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

The dose response effects of injected polynucleotide in muscle *in vivo* is determined as follows. Suitable template DNA for production of mRNA coding for the polypeptide of the present invention is prepared in accordance with a standard recombinant DNA methodology. The template DNA, which may be either circular or linear, is either used as naked DNA or complexed with

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liposomes. The quadriceps muscles of mice are then injected with various amounts of the template DNA.

Five to six week old female and male Balb/C mice are anesthetized by intraperitoneal injection with 0.3 ml of 2.5% Avertin. A 1.5 cm incision is made on the anterior thigh, and the quadriceps muscle is directly visualized. The template DNA is injected in 0.1 ml of carrier in a 1 cc syringe through a 27 gauge needle over one minute, approximately 0.5 cm from the distal insertion site of the muscle into the knee and about 0.2 cm deep. A suture is placed over the injection site for future localization, and the skin is closed with stainless steel clips.

After an appropriate incubation time (e.g., 7 days) muscle extracts are prepared by excising the entire quadriceps. Every fifth 15 um cross-section of the individual quadriceps muscles is histochemically stained for protein expression. A time course for protein expression may be done in a similar fashion except that quadriceps from different mice are harvested at different times. Persistence of DNA in muscle following injection may be determined by Southern blot analysis after preparing total cellular DNA and HIRT supernatants from injected and control mice. The results of the above experimentation in mice can be use to extrapolate proper dosages and other treatment parameters in humans and other animals using naked DNA of the present invention.

It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference.

Sequence Listing

	(1) GENERAL INFORMATION:
5	(i) APPLICANT: Human Genome Sciences, Inc., et al.
	(ii) TITLE OF INVENTION: 207 Human Secreted Proteins
10	(iii) NUMBER OF SEQUENCES: 800
15	(iv) CORRESPONDENCE ADDRESS:
10	(A) ADDRESSEE: Human Genome Sciences, Inc.
	(B) STREET: 9410 Key West Avenue
20	(C) CITY: Rockville
	(D) STATE: Maryland
25	(E) COUNTRY: USA
23	(F) ZIP: 20850
30	(v) COMPUTER READABLE FORM:
	(A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
35	(B) COMPUTER: HP Vectra 486/33
33	(C) OPERATING SYSTEM: MSDOS version 6.2
	(D) SOFTWARE: ASCII Text
40	
• •	(vi) CURRENT APPLICATION DATA:
45	(A) APPLICATION NUMBER:
43	(B) FILING DATE:
	(C) CLASSIFICATION:
50	
	(vii) PRIOR APPLICATION DATA:
55	(A) APPLICATION NUMBER:
	(B) FILING DATE:

	(viii) ATTORNEY/AGENT INFORMATION:	
5	(A) NAME: Kenley K. Hoover	
	(B) REGISTRATION NUMBER: 40,302	
	(C) REFERENCE/DOCKET NUMBER: PZ007PCT	
10	•	
	(vi) TELECOMMUNICATION INFORMATION:	
15	(A) TELEPHONE: (301) 309-8504	
13	(B) TELEFAX: (301) 309-8439	
20		
20	(2) INFORMATION FOR SEQ ID NO: 1:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 733 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:	
	GGGATCCGGA GCCCAAATCT TCTGACAAAA CTCACACATG CCCACCGTGC CCAGCACCTG	60
	AATTCGAGGG TGCACCGTCA GTCTTCCTCT TCCCCCCAAA ACCCAAGGAC ACCCTCATGA	120
35	TCTCCCGGAC TCCTGAGGTC ACATGCGTGG TGGTGGACGT AAGCCACGAA GACCCTGAGG	180
	TCAAGTTCAA CTGGTACGTG GACGGCGTGG AGGTGCATAA TGCCAAGACA AAGCCGCGGG	240
40	AGGAGCAGTA CAACAGCACG TACCGTGTGG TCAGCGTCCT CACCGTCCTG CACCAGGACT	300
	GGCTGAATGG CAAGGAGTAC AAGTGCAAGG TCTCCAACAA AGCCCTCCCA ACCCCCATCG	360
•	AGAAAACCAT CTCCAAAGCC AAAGGGCAGC CCCGAGAACC ACAGGTGTAC ACCCTGCCCC	420
45	CATCCCGGGA TGAGCTGACC AAGAACCAGG TCAGCCTGAC CTGCCTGGTC AAAGGCTTCT	480
	ATCCAAGCGA CATCGCCGTG GAGTGGGAGA GCAATGGGCA GCCGGAGAAC AACTACAAGA	540
50	CCACGCCTCC CGTGCTGGAC TCCGACGGCT CCTTCTTCCT CTACAGCAAG CTCACCGTGG	600
	ACAAGAGCAG GTGGCAGCAG GGGAACGTCT TCTCATGCTC CGTGATGCAT GAGGCTCTGC	660
	ACAACCACTA CACGCAGAAG AGCCTCTCCC TGTCTCCGGG TAAATGAGTG CGACGGCCGC	720
55	GACTCTAGAG GAT	733

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	(A) LENGTH: 5 amino acids	
_	(B) TYPE: amino acid	
5	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:	
	Trp Ser Xaa Trp Ser	
10	1 5	
15	(2) INFORMATION FOR SEQ ID NO: 3:	
13	(2) In diameter to the one of	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 86 base pairs (B) TYPE: nucleic acid	
20	(C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:	
25	GCGCCTCGAG ATTTCCCCGA AATCTAGATT TCCCCGAAAT GATTTCCCCG AAATGATTTC	60
4 3	GCGCCTCGAG ATTICCCCGA AATCIAGATT TCCCCGAAAT GATTICCCGG TUTTO	
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35	(A) LENGTH: 27 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(D) TOPOLAGI: IIIleal	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:	
	GCGGCAAGCT TTTTGCAAAG CCTAGGC	27
45		
	(2) INFORMATION FOR SEQ ID NO: 5:	
50	(i) SEQUENCE CHARACTERISTICS:	
50	(A) LENGTH: 271 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:	
J.J	•	60
	CTCGAGATTT CCCCGAAATC TAGATTTCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG	60
۷0	AAATATCTGC CATCTCAATT AGTCAGCAAC CATAGTCCCG CCCCTAACTC CGCCCATCCC	120
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	GCCCCTAACT CCGCCCAGTT CCGCCCATTC TCCGCCCCAT GGCTGACTAA TITTTTTTAT	180
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	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:	
35	GCGAAGCTTC GCGACTCCCC GGATCCGCCT C	31
40	(2) INFORMATION FOR SEQ ID NO: 8:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 12 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:	
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55	(2) INFORMATION FOR SEQ ID NO: 9:	
	(i) SEQUENCE CHARACTERISTICS:	
60	(A) LENGTH: 73 base pairs (B) TYPE: nucleic acid	

	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	•
_	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:	
5	GCGGCCTCGA GGGGACTTTC CCGGGGACTT TCCCGGGACT TTCCATCCTG	60
	CCATCTCAAT TAG	73
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	(2) INFORMATION FOR SEO ID NO: 10:	
. ~	(a)	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 256 base pairs	
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	(D) TOPOLOGY: linear	
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	CTCGAGGGGA CTTTCCCGGG GACTTTCCGG GGACTTTCCA TCTGCCATCT	60
25	CANTTAGTCA GCAACCATAG TCCCGCCCCT AACTCCGCCC ATCCCGCCCC TAACTCCGCC	120
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30	GGCCGCCTCG GCCTCTGAGC TATTCCAGAA GTAGTGAGGA GGCTTTTTTG GAGGCCTAGG	240
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	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
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	GACAGGCTAT COGAGAATCT GAGAGCTGGG CCCGGCAATT CCTCCAGYTA CCCTTGTGAC	60
	CTAAGTCCAG TCACACATTT CCCAAAGTTT CTCTTTGTCA TAACCCTGGT CTGGCTGGTT	. 120
50	TTGRGGRCTT GAGAATGGGT CAGGGACTCC AGGCCAAGTC CAACAGAGAC CCCAAACCCA	180
	CCACACACCA GCAGCCACAA CCTCACCACC AACAAAGAGG ACTTTTGTGG GGCCACAAGT	240
55	AAGAGGTCAT TTCTGGAATG GACTCAGACC TTTAAACAGG AGAGTTGAGC ACTTCCAGKS	300
-	AGTITTIAAG CAAGGCATGG GGAACAGGGA ATAGAACCTT TCAAAGAGGT TGCCCAGAGA	360
	AAAGCTGGGC CTCTTGCATT CGGCTTCCTT GGAGCAGCCT CTTCTGGCAG AAAGCCATCA	42
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	GCCCCGTGCC CTGAGGTGAC TTCCTAACTA TGTGGTTTCA TTAGCGAATT TATTTTTTGT	1020
20	CCTCGGTGGA CATTTGTATT TTGTTAGGTT GCTGTTTAAG CTCAAGTTTG CTGTGCTCTC	1080
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40	GAAAAGCCAC GTCATATAAC TCAAGAATAA ATGGTGTTTT GGAAATTTTA AAATTATCAT	1680
	CGAAGGTGGT GAAACTATTT CAGGCCCAAA TGAAAGGAAA TCGCCAGTTG GGGATGAAAT	1740
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45	CATTITIANA AGIGCGCATG ATTCTACATA TGAGAATTCT TTAGGCCAAG ANACTGTCCT	1860
	TOSCTCAGAG GTGTTGGGAA TTAAAGCAGA GAGAAGCCAT TCGTGATGCT TAGAACCAAG	1920
50	C10771 CON	1980
	GARAGCACTT TGTAGGGGAA CTTTAGTAAG TTCTTCTCAT TTCATTATGT TTCTTCCAAG	2040
	GAAACAGGAG AGACTGAATT AATAATTCTC TCTTTCCTCT TAAGCACTTT TAAAATAATA	2100
55	AAGTACATCT TGAAATTTGG GGGGCCATCT CTGATTTAAA AAAAGAAAAA GGCTGCTTGA	2160
	TGTATGTTAT GCAGAGACAC TCTGCCTCTG GTGGCTGCAG AGCAATACCC AAGCCTCATT	2220
60	The second secon	228

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	ACGGGATGGT GGGTCTGGGA CCCCAATTCA TTCTTATCTG CCAAAGAATT ATCTAGAAGC	2340
_	ACATCAAATA CCAGCACCCC ACCTGCACAA TGGGGGTGGA AAACTTTTGT ATCCCTAAGC	2400
5	ATATTATTTT ATACTCTCTC CCATCCCATC TGGAAATACT TTATTTTTAA CCTCAGGATT	2460
	TARATARAGT ARACACTATG ACATITARAR ARARARARA ARACTCGAG GGGGGCCCGG	2520
10	TACCCA	2526
15	(2) INFORMATION FOR SEQ ID NO: 12:	
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:	
25	CACTGCACCA GCTTTGTTAT CTGTAAAATG ATGATAATAC CAACACCTTC TTCTTGGGGT	60
	ACTGAAGATG AGAGAACATG ATATGTGTAA AGTGCCTTCC ACAATACCCA GAACATAGCA	120
20	AACATGTAAT GAATGTAGTA ATAGTAATTA TTTTATTTTC TTTTGATTCA GTTGGGACTA	180
30	TGTTCAGCTG TAACAGAATA CCCAAAATAA CTGTTTTAAA CAAATTAAAG TTTWGTTGTG	240
	AAGTTTIGTT ACGAATTCAG ACAATCCAGG GCTTTTATAG ATGCACCAGG ATCAGCAGGT	300
35	ACAAAGGCAT CTTTCCTGAT TTCTGCCAGT CTCAATGCAT GGGTTGCAAT CCAGARTCCA	360
	RGATGGCAGT TOCAGCCCTG GTTACGCCCA TATTAGCACA CAGAAAGAAA GAGAAAGGGA	420
40	TGTGCCTCTT CACTTTAATC ATAGCTCCCA CTAGATGCAC CCACTACTTC TGCTGATACT	480
40	CCATTAGCTA ATGCTTGCTT ACATGGTCAC ACTTAGTTTC CAGAGAGACA TGTCTGGACA	540
	GICATGIGCT CAATTAATAT CCAAGIGTCC AATTACTGAG AAAAAAAGAA ACTAGCACCT	600
45	TIGGITGGIT GCATICCICT TAGCATAAGC CACATICTIT TIATGAAGIT GICCICAGIT	660
	ACTTOGATIC CTCAGTTGTC CTTTCAWTTA GAAAWGCYCC TKGGACAYCC TGAAWCTGAC	720
	TTCTTTTGTC ATCAGCACCA TCACTACCAC TGCCYTCTTC AAAGCCACCA CGTTCTGTCC	780
50	CCAGGATGGT TGCAACAACC ACCATAGGGA CTTTTTGCCT TCTACTTCCA CACAATAGNC	840
	CAGAGTAAGC TTTTGAAAAT GTAGGTCAGA TCATGTCTCT CTCTTCCTCT TCAAAACCCT	900
55	CCCGATGGCT TITCATATTA CTCAAAAGAA AACCTAAAAC TITGCTGTGA GATCTATGTG	960

ACCOGNITA TICTICCICT TACTITATCT CTGTATTGCT CTTCCTCACT CTACTCCAGC

CATCCCACCT CCTTGCTGCT TGTCCTATAC TCCTAAAAGA AGTTCAGTCT TCCCTTATGA

272

	TATTTGCACT TAAAATAGAA AAAAAAAAA AAAAAAAACT CGAGGGGGC C	1131
		-
5	(2) INFORMATION FOR SEQ ID NO: 13:	
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 941 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:	
13	GGCACGAGTA GCATTTCATT TAATCTGCAG GTATATTCTC CCAACAGTTT ATTGTCATGT	60
	GATGTCCTCA GCCAAGATTG TRAGGCAGAG AGGAGCTGTC CCAACCTACT ATACCACCGA	120
20	GGCTGGAGAG ATCATATTTT TGGTATTAAA CTGGAGTCTC TCCATCCTTC ACATTGTTGA	180
	TGTCCTCTGT AGCAAACCGG AAAAGTCAGT GACAGAAGAT GCCGCTAGCG GTTTGAGCCA	240
25	GAGAATGACA GCTCTGGTTT GGAGAAAAGG GCCGGATGGT GGCTCTAGAA AGCCCATCCT	300
23	TCTGCTCTTC TTTTTTCTCC CCCTTATATT GTGCTTTCAT TCATTCATTC ATTCATCAAA	360
	CATTIGITGA GCACCIATTA TGTGTCAAGC TCTGTGCTAG CCTCTGGAAA ACCTGCCCTC	420
30	ATGTAGCTCA CTGTGGAGTA GGAGAAACAA TGACTACACT ATGATAAGCA CGGGTTGTCA	480
	GGGTCTCACA GAGCAGTGGC CCCTCATCCA GACCGATGAG GTCAAAGAAG GCATCCAGGC	540
35	GAGGATGGTG TCAGAGCTAA CTGAAGAATG AGAGGGAGCT GCACCASCAG GGGTTGGAAC	600
33	TGAAGGTGGC AGTGCCTGGA GTCTTGATTC CAGCAGAGGG AGAGCAGTCT GTGAAAAGGC	660
	ACCAAGGGTG GGAGAGGGCA GAGCACATGG AGGAACTTCA GGTAGTTCTG GATGGCSCTG	720
40	GGGCAAAGCT AGAGAGGTAA GAAGAATCTA CAAATGTTCC TCGAGTTACA TGAACTTCCA	780
٠.	TCCCAATAAA CCCATTGGAA ACGAAAAATT TAAGTCAGAA GTGCATTTAA GGCTGGTCCG	840
45	AGTAGAATGA TITTTACAAC GAATTGATCA CAACCAGTTA CAGATGTCTT TGTTCCTTCT	900
43	CCACTCCCAC TGCTTCACCT GACTAGCCTT TAAAAAAAAA A	941
50	(2) INFORMATION FOR SEQ ID NO: 14:	
	(i) SEQUENCE CHARACTERISTICS:	
55	(A) LENGTH: 843 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

(D) TOPOLOGY: linear

	CNAGGGATAA CCCCAAAGNT GGGAAATAAA CCCTCAATTA AAGGGGGAAC CAAAAAGCTG	6Ò
	GGAAGTTCCC CCCCGCGGTG GCGGCCNGNT CTAGGAACTA GTGGAATCCC CCGGGGCTGC	120
5	AGGGAATTCG GCACGGAGTG GGAATGTTGT TTGTATGATA CTATTTCCAC AAWATGCATT	180
	GAGACTTGGT KTGTGGCCTA GGACATGGTC AATTCTTTYT AAATATTCCG TGAATTTCTT	240
10	TAGTICCATAT TCTCCGATGG GGCCTGTGGG GACAGAGTTC TAAATATGCC CATTAGATTA	300
10	AATCTCTTCA TTCTGTTGCT CACATCTTCT ATATCCTTAT TAATCTGTCA ATCTCTTCAA	360
	GAGAGGTGTT ATTAAAATCT CTCACTGTAT GTGTCACTTT GCCCTTAAAA TTCTGATGAT	420
15	TTGCTTTATA AATGGTTATA ACCATTTCC AGGAAGAACA TTAAAGAACT TTCCATTGGC	480
	ATTATCCAGT TICCCTCAAA ATACTGGTTT TTTTTATTTT GGCTNCTAAG CAGCTATGAA	540
20	TCCAGTTTCT CAGAAGCCCT TGTCTCAAGG CATTTGTTTC CAGATTACCT TGTTAGCATC	500
20	CACACTATGG GCTATTTTAG AAAAACAAAA AAAGTATCAA AATCATATAG CTATGATTTT	560
	CCTGTGCTTG AAGGAGCCTT AAAGCTCATC TAGTCCAGCC AGTATTTGTT CATCCAAATT	720
25	CTGCCAAGAA ATCTCTATTG TCAAGATATT CTTTACCATC TTTGGGACAT TCTCATTATT	780
	AGAAACAAAT CCTAAGAAGA AATTCTGCCA TAKACAACCC ATCCGTTCTT TAAAAAAAAAA	840
20	AAA	343
30		
	(2) INFORMATION FOR SEQ ID NO: 15:	
35		
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1018 base pairs (B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:	
٠.	CIGIAATITI TAATITICAT ATACCGIGCT TIGATICTAA TITTATITIT TGAGTICTCT	60
45	GAAGGITACA TATACAGAGT GCTICAGGAA TGATCATTIT GTTATTATTC ATGCTTCTTA	120
	ACAATGTTGT TTTAGTCCAA GAAGATAATT GCCAGAGAAA GAATACAGTG CAGGAAAGAA	180
50	GARGCTGGAG CCAGTGGTGA AGARGATTG AGARGACAGA CATTGTGGGA ATGAAATCAT	240
50		300
	GAATAATCGT GTTTTTGAAT TGTCCAAAAA CTTCTACAAA CCATGAAATG TTGGAGTTTA	
5 5	AATCTAATTG TTGAAAAATT CCCCACATTC CTTGTATCCC TTAGGTTGAG CATAATTCCA	360
	CATCOGTOGA CTGATGCACT TCCCAAGAGG GGGCCTCATT AACTCTTCCG AGGCAGCAGC AGCAAGGGCA CCCCCTCCTT TCCCCCCACA CCCCAYTTCT CATGGCTCTT CTTTCTCTCA	420 480
	successions condition incomme admitted and an entitle and an entit	-200

60 TCTCATGCTT AGGTTAGAAA AGGGCACAAG GTAAGGAAGC CCTTGGGAAT AGGCTGAATC

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	TGGCTATCTA ATTTGGTGCC AAATACTTAA TGTGCTTGAA TTTAAAAACA GCAAACATGT	600
	AGAAAGGTAA TTATAATTAT GAGGCCAGTT CTTTAAGCTA GCTTTTTTTC CCCTCTCAAA	660
5	CAGCATATTG GCTTGGATGT CAGCAGGAGA AAGTGTTTTT TGCAATACAC ATAATGCATA	720
	TATGGTCCTG TTAGCAATCT ATAGAAAATA GATATTGCTC ATTAAGGTAA ATATTTTTGT	780
10	TGATGAATGA TCTGGAATGG TCTGGACTTG TTGTGTGAAC AGGAAATTGC TCTGTAGGCT	840
	TTGACTTGTG AGGTAAAGAG TGAGGCTGGT AAGATTAATT AAAGTAAATA CTGTGACAAT	900
٠, ٣	AGGATGTCAA AACCAAAAAC GTGTTTCTGA AACTCAAGGA ATTAATGACA CATAGGGAAG	960
15	TTTTTGCCAT ATTAAGCATA GAGTAGGAGA GGCAAGTCAA GAATAAAAAA AAAAAAAA	1018
20	(2) INFORMATION FOR SEQ ID NO: 16:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 661 base pairs	
25	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

(D) TOPOLOGY: linear

TTTAAGAAAT TAGTGAATCC CCGGNTGCAG GGAATTCGGC ACGAGGAGGA GGCCGTCAGC 60 TGGCAGGAGC GCAGGATGGC AGCTGYTCCC CCGGGTTGCA CCCCCCCAGY TCTGCTGGAC 120 ATAAGYTGGT TAACAGAGAG CCTGGGAGCT GGGCAGCCTG TACCTGTGGA GTGCCGGCAC 180 CGCCTGGAGG TGGCTGGGCC AAGQAAGGGG CCTCTGAGCC CAGCATGGAT GCCTGCCTAT 240 GCCTGCCAGC GCCCTACGCC CCTCACACAC CACAACACTG GCCTMTCCGA GCTGCTGGAG 300 CATGGAGTGT GTGAGGAGGT GGAGAGAGTT CGGCGCTCAG AGAGGTACCA GACCATGAAG GTGCGCAGGG CAGGGCTCGG ACCTACCCCA GGAATGTCCT GCCCTGGGAA TGACAACACA 420 GTCCACACCA TGCACGGGGA GGCAAACAGG GGCAGCTGAC CCAGCCCAGG GGTCAGANGA 480 45 GGTCTTGCCG AGGAAGTGGC AGCTAAGCTG ATACCTGATA TGCACWAGKC AGCCARGYGG 540 AGACAGGCAA GGAAGAAGCT TGTTTTGAGG ACAGAATTTT CTAGATCACT CAGCACCATC 600 50 TOCCTTTTGG GGCTTTTTGT TTTATTTTGT TTTTGAGACG GGGTCTCGCT CTGTCGCCCA 660 661 N

55

(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS: 60

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	(A) LENGTH: 553 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	·
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:	
	GGCACAGGGC TATTTGCCCC TCTCTCCACA TGACAGAACT GCTCTAAGTT TCTTTGCTGC	60
10	TCTTCTCAGC TGTCAGACGG CTTGCTGCTT GTTTTCCACA CCACCATGTC TATTCTTTGC	120
	TGTCCTTWAC TCTGCCTGTT TTTTTCCTTT TGTATTTCTT CTGGCTCTTG TCCCTTTTCC	180
	CACGTGTCWC AGCTTTCCTT TATTGCCACT TICAGTCAGA GCAGTCCTGT GCTTCTGGTG	240
15	CCGGCATACA ATACTTACTT GAGTTTCTTG GCTTTTCTTG ACTGTGCATC TCTTACTTCA	300
	ACATAGGAAT AGCCTGTCAT AGAATTTCTC CAGTTCCAGG GCTCAAGAGG GAGAGTGCCA	360
20	GAAAATTGAG ACTGTTTTCC CTGTCTTGGA TTGAATTCAT AAAGCAAAAC CAGTGTTTGT	420
	GTGAGGGTTT GCTGTGTCAT GCCTATAGGT TGTTTGGGTG CAAACCTATA GAATCCAGCC	480
	TGCGAAAAGA AAGRAACCAG AGAATANCAG CATCAGAACA ATGCTTGACA TCATTTCTCA	540
25	ATCAAGCAGT CCA	553
30	(2) INFORMATION FOR SEQ ID NO: 18:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 869 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:	
40	GGCACGAGCT GCCAACACTG AGGTCTTCGT GGCTTCTCAC ATCTAGATGT ATCCCTCTCA	60
	AATCTATCCT CTATCCAGGC ACCAGATTGA GGTATCTAAA ATGTCAACTT TCCAGTTACT	120
45	CCTTCTTATA CTAGCCCAAT CAACTTACAA GATAAAGTCC AAGCCCCTTC ATATGACAAA	180
	CCACACCCTG CTTAACTCTC CAGGITTGAA TCCTTCATCT CCTACTTTAA ACTITAAAAC	240
	CCAGCAGCAC GAAAGTGTCT CCTATGCATG TTGCCATATG CGTTCTCTCC ATCATGCATT	30
50	TOCCTGAGCA AGATGTCTTG AGTTAACATC TTATTCTTTA AGACTCATTG TGGTGGTAGA	36
	CAGCCTTTAA TAACGGATCC TTGGCCAGGC ACAGTGACTC ACACCTGTAA TCCCAGAACT	42
55	TTGAAAGGCC AAAGAAGGAA GAAAGCTTGA GGCCAGTAGT TTGAGACCAG CCTGGGAAAC	48

AGAGAGATAT CCCATCTGTA CCAAAAATTT AAAAAAATAT TAGCAGGGAG TAGTGGCATG
CACAAGTGGT CCCAGCTCCA TGGGAGASTG AGGTAGGAAC ATCACTTGAG CCCAGGAAGT

	CAAGGCTGCA GTGAACCATG ATCAGAACAT TGCANTCCAG CTTGGGTAAC AGAGTGAGAC	660
	CTTAGGTCAG AAAAATGAAT AAATAAGCAT AAAATTITAA AAACTTAGCC AGGCATGGTG	720
5	GCACACATCT GTGGTCCCTG CTACTTAGGA GGCTGAGGTG AGAGGATCCT TGAGCCCAGG	780
	AGGICAACAC TACAGIGAGC TATGATIGIG CCACTAAACT CCAACCIGGG IGAAAAAGCA	840
	AAACCCTGCC AAAAAAAAAA AAAAAAACT	. 869
10	•	
	(2) INFORMATION FOR SEQ ID NO: 19:	
15		
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 959 base pairs	
	(B) TYPE: nucleic acid	
••	(C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:	
25	GGCGAGCCGA GATCGTGCCA TTGCACTCCA GCCTGGGCAA CAAGAGTGAA ACTCTGTCTC	60
23	AAAAAAAAA AATTATAATA CTATATGCCA TAAAATGACA TTTCATATTT AAAGAGTTTT	120
	TTAAAACTCT TGTATTCACA TGCCATAATT TGAAACCCTA TTTCACTGAA TGAGAATGGT	180
30	ATCTGTTGTC CTCATTTTTT CATTTTTATC CTTAACAATT TCCACCACAG CCAGTGCATA	240
	TAATGGCAAT GACACCCAGG GATGGAATGA TAAGTTCCAT CRCMGCTCAG TCAAGACGCA	300
35	GACTTGATGT GGCCCCAACA ACAGTCAATA ATGGAGTCTC CAAAATAAAG CTCTATAGGA	360
J J	AAGGTAAATA CCCGCTGCAC AAGAAACCAC AGCATCTAGG TTCTAACCCC ATCTCTATGA	420
	AGAGCTTGCT GGGAGAGTTT TGACATTWAA CAATCTGTCT GATKGCCAAT TTTYTTCTTC	480
40	TATAAAATGA TAATGITKGA YTCAAAGATC CAAAGTCAAT TCATGGTCTA AAACTTAATG	540
	ATTITITAG GITTIGKGAC ATTICACTOT ACACTOTAGI AATTIATATC TIATTITCCC	600
45	ACTAATTTAG AAAAATATYT AAATGATCCT TAATTGGCAA TGGGTCCTAA GAATTTTGTT	660
	TTAAATCCCT GTTACCCAAA AGAGCCCTTT TTTGTATCTC GCAGTAGTTA CAAGGATCTT	720
	TCTAAATCTT AAAAAAAAA AAAAAAGAAA GAAAGAAAAG AAAAGAAAAA AAGTCAGCCG	780
50	GGCGTGGTGG CTCATGCCTG TAATCCCAGC ACTTTGGGAC CAAGGTGGAC AGATCACGAG	900
	GTCAGGAGAT GGAGACCATC CCGGCCAACA TGGAGAAACC CTGTCTCTAC TAAAAAAAAAA	959
55	AAAAACTCGA GGGGGCCCG GTACCCAATN CGCCGGCTAG TGGTCGTAAA ACAATCAAA	737

⁽²⁾ INFORMATION FOR SEQ ID NO: 20:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1446 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

10	COGGGCAGGG CTGTGTGGCA CCGCCAGGGA GCGGGCCCAC CTGAGTCACT TTATTGGGTT	- 60
10	CAGTCAACAC TTTCTTGCTC CCTGTTTTCT CTTCTGTGGG ATGATCTCAG ATGCAGGGGC	120
	TOGTTTTGGG GTTTTCCTGC TTGTGCCAAG GGCTGGACAC TGCTGGGGGG CTGGAAAGCC	180
15	CCTCCCTTCC TGTCCTTCTG TGGCCTCCAT CCCCTCATGG GTGCTGCCAT CCTTCCTGGA	240
	GAGAGGGAGG TGAAAGCTGG TGTGAGCCCA GTGGGTTCCC GCCCACTCAC CCAGGAGCTG	300
20	GCTGGGCCAG GACCGGGAGA GGGAGCACTG CTGCCCTCCT GGCCCTGCTC CTTCCGCAGT	360
20	TAGGGGTGGA CCGAGCCTCG CTTTCCCCAC TGTTCTGGAG GGAAGGGGAA GGAGGGGGTC	420
	TTCAGGCTGG AGCCAGGCTG GGGGTGCTGG GTGGAGAGAT GAGATTTAGG GGGTGCCTCA	480
25	TOGGGTOGGC AGGCCTOGGG TGAAATRAGA AAGGCCCAGA ACGTGCAGGT CTGCGGAGGG	540
	GAAGTGTCCT GAGTGAAGGA GGGGACCCCC ATCCTGGGGG ATGCTGGGAG TGAGTGAGTG	600
30	AGATGGCTGA GTGAGGGTTA TGGGGAGCCT GAGGTTTTAT GGGCCTGTGT ATCCCCTTCT	660
•	CCCGGCCCCA GCCTGCCTCC CTCCTGCCCG CCTGGCCCAC AGGTCTCCCT CTGGTCCCTG	720
	TCCCTCTGGT GGTTGGGGAT GGAGCGGCAG CAAGGGGTGT AATGGGGCTG GGTTCTGTCT	780
35	TCTACAGGCC ACCCCGAGGT CCTCAGTGGT TGCCTGGGGA GCCGGACGGG GCTCCTGAGG	840
	GGTACAGGTT GGGTGGGCCC TCCCTGAGGG TCTGGGGTCA GGCTTTGGCT CTGCTGCCTC	900
40	TCAGTCACCA AGTCACCTCC CTCTGAAAAT CCAGTCCCTT CTTTGGATGT CCTTGTGAGT	960
	CACTCTGGGC CTGGCTGTCG TCCCTCCTCA GCTTCTTGTT CCTGGGACAA GGGTCAAGCC	1020
• .	AGGATGGCC CAGGCCTGGG ATCCCCCACC CCAGGACCCC CAGGCCCCCT CCCCTGCTGC	1080
45	TTTGCGGGG GCAGGCAGA AATGGACTCC TTTTGGGTCC CCGAGGTGGG GTCCCCTCCC	1140
	AGCCCTGCAT CCTCCGTGCC STAGACCTGC TCCCCAGAGG AGGGGCCTTG ACCCACAGGA	1200
50	COTOTOGTOG COCCTOGCAC TCAGOGACCC CCAGCTCCCC CAGCCCTGGT CTCTGGCGCA	1260
50	TCTCTTCCCT CTTGTCCCGA AGATCTGCGC CTCTAGTGCC TTTTGAGGGG TTCCCATCAT	1320
	CCCTCCCTGA TATTGTATTG AAAATATTAT GCACACTGTT CATGCTTCTA CTAATCAATA	1380
55	AACGCTTTAT TTAAAGCCAA AAAAAAAAA AAAAAACTCG AGGGGGGCC CGTACCCAAT	1440
	TCGCCA	1446

(2) II	VEORMATION	FOR	SEQ	ID	NO:	21:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1471 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:	
	CAAAAAATAA TAATGATAAT TTAAAATAAA TAAGTAACTA ATAAAAAGAT TTTATATCCC	60
15	AGTOTTATGA TGTTGGTTGG CAAGGCTAGA TAAAAAGATG TTAGAATGAA AGAACATATT	120
	TTTAGTGATA TGTAAATGAA GGATTCTACA ATAGTCATAT ATTTTTATAT GAATGAATGT	180
	TOGGTTOGGC TGGAGAGGTA TGTGTGTGTA AATATAAAGG TCTCACATTC AGAGTATAGC	240
20	TCTGAAATAA TGGAACTCAT GTCTACAATT CAACATGCAT CTGTATAGTT ACATCTCATG	300
	TAAATATACA CAGACATATT TTGCAGCCAG TAATTGACAG TTAATGTCCA AAACAGGTGA	360
~ =	TTGATAGGTA ACAGAAATTA GATAACCACC AATTTTGCCC AAGAGAAAGA CTAGAAGGAC	420
25	TARARGCAGT TGRATGTATG GTACTGACAT TGTCATARGC AGTCTGATAR CCAGTTTATT	480
	GAAACGTGTG CATTAACAGA GAATTTAATT TTAAACCCAT AATTTCTCCT ATCCATTAAA	540
30	ATATTATAAT TGTTAGTAGT ATGAAACCAA CAGGAAATGT TTTTTAATCA TTTAGTGAGG	600
	TGATTCATTT GTTTCATGGG CAAACACTAT CCAGGAAAAG CCTTGCTTGC CTGTTTCCCA	660
0.5	AAGAGCTCTA AGAAATAGAA TCAAGTGTAA AATGGTTCAG ACCATTCAGG ATTTCTTGTC	720
35	ACTOTTOTCA ACCOCGATOT TOCTGTTATT ACTGATGTTT GAAACCCTGT CATTAGCCCC	780
	GCCTGGTTA AAGCCCCTCA GAGTCACCTC TCATTCATAG CAATAGAATT CAACCCCAAG	840
40	TEGTTGATEG TETCCCCAGC ACAGCCGAGA GACCTGATCT CTGGATTCAG TECTTTTAGC	900
	TCTTCGAGTT TACCCTAAGA TACCTTCGGG CAATATTTTT AACCAACCCA AAAGCTCTTC	960
	AGGTCATTTC TGAAGAGGAC AAGGTGAATC TTGGCTTGGA ACACCATTTT TGGGCTCTTG	1020
45	CTACTGAATG AATCAGAAAG GAATTTTTC TGAAGAGCAT TAGAAAGTAA AGGAGATGTT	1080
	AAAATAAGIT CITGAAGTAT GITTTATATT TATCTAAAAC ACTGATTITA AAAGITTACA	1140
50	TICAAATGIG TATICAAAAG AAGTACIGAT TIGIAATTAT TATAGITIGI GIGTATCATC	1200
	CCCTTTTAAC CGIGCCTAAC AACTGTACTI AAATTTTGTT TICCTAGTGT AACAAATGTT	1260
	TCCCATAAGA TTTTCTAGAG CCAAATAATG GGAGTGAAAA ATTCCTTAAG TGTTATATAA	132
55	GAAAATATAT TAGAAAATCA GCTTTGGATT ATACGATTTC TAAAATATAC TAATACAGAA	138
	TCCTCAGTAA TATGTTTTGA ATTGGATTTT TTCTCAGAAC TGTTACATAA TAAATAATAC	144
60	ATCAACCAGA AAAAAAAAA AAAAAAATTN C	147

5	121	INFORMATION	FOR	SEO	IĐ	NO:	22:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1402 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

	(XI) Sugarian believe to the termination of the ter	
15	AGGGACGTCT TGCCTGAGGA GATGCCCATT TCTGTCCTGG RTTACCCTCA CTGCGTGGTG	60
	CATGAGCTGC CAGAGCTGAC GGCGGAGAGT TTGGAAGCAG GTGACAGTAA CCAATTTTGC	120
20	TOGAGGAACC TCTTTTCTTG TATCAATCTG CTTCGGATCT TGAACAAGCT GACAAAGTGG	180
20	AAGCATTCAA GGACAATGAT GCTGGTGGTG TTCAAGTCAG CCCCCATCTT GAAGCGGGCC	240
	CTAAAGGTGA AACAAGCCAT GATGCAGCTC TATGTGCTGA AGCTGCTCAA GGTACAGACC	300
25	AAATACTTGG GGCGGCAGTG GCGAAAGAGC AACATGAAGA CCATGTCTGC CATCTACCAG	360
	AAGGTGCGGC ATCGGCTGAA CGACGACTGG GCATACGGCA ATGATCTTGA TGCCCGGCCT	420
30	TGGGACTTCC AGGCAGAGGA GTGTGCCCTT CGTGCCAACA TTGAACGCTT CAACGCCCGG	480
30	CGCTATGACC GGGCCCACAG CAACCCTGAC TTCCTGCCAG TGGACAACTG CCTGCAGAGT	540
	GTCCTGGGCC AACGGGTGGA CCTCCCTGAG GACTTTCAGA TGAACTATGA CCTCTGGTTA	600
35	GAAAGGGAGG TCTTCTCCAA GCCCATTTCC TGGGAAGAGC TGCTGCAGTG AGGCTGTTGG	660
	TTAGGGGACT GAAATGGAGA GAAAAGATGA TCTGAAGGTA CCTGTGGGAC TGTCCTAGTT	720
40	CATTGCTGCA GTGCTCCCAT CCCCCACCAG GTGGCAGCAC AGCCCCACTG TGTCTTCCGC	780
40	AGTCTGTCCT GGGCTTGGGT GAGCCCAGCT TGACCTCCCC TTGGTTCCCA GGGTCCTGCT	840
٠.	COGAAGCAGT CATCTCTGCC TGAGATCCAT TCTTCCTTTA MITCCCCCAM CCTCCTCTCT	900
45	TOGATATOGT TOGTTTTGGC TCATTTCACA ATCAGCCCAA GGYTGGGAAA GCTGGAATGG	960
	GATGGGAACC CCTCCGCCGT GCATCTRAAT TTCAGGGGTC ATGCTGATGC CTCTCGAGAC	1020
50	ATACAAATCC TTGCCTTTGT CAGCTTGCAA AGGAGGAGAG TTTAGGATTA GGGCCAGGGC	1080
30	CAGAAAGTCG GTATCTTGGT TGTGCTCTGG GGTGGGGGTG GGGTGTTTCT GATGTTATTC	1140
	CAGCCTCCTG CTACATTATA TCCAGAAGTA ATTGCGGAGG CTCCTTCAGC TGCCTCAGCA	1200
55	CTTTGATTTT GGACAGGGAC AAGGTAGGAA GAGAAGCTTC CCTTAACCAG AGGGGCCATT	1260
·	TTTCCTTTTG GCTTTCGAGG GCCTGTAAAT ATCTATATAT AATTCTGTGT GTATTCTGTG	1320
۲۵	TCATGTTGGG GTTTTTAATG TGATTGTGTA TTCTGTTTAC ATTAAAAAGA AGCAAAAATA	1380
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(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1047 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

15 GCCACAGGGG ACTACAGGCA CCCACGACCA TACCCAGCTA ATTTTTGTAT TTTTTTGTAG 60 AGATGGGGTT TCACGATGTC GCCCAGGCTG GTCTTGAACT CCTGGGCTTG AGCGATCTTC 120 CCATCTTTCC ATCTTGGCCT CCTAAAGTGC TGGGACTGCA GGCATGAGCC ACCATGCCCA 180 20 GCCAAGATTC TTATTGATTA CCATGTTGCT TCAAGAAGCC AAGCCAGTTT CCAATATTCC 240 CCATTTCCTG GAGTCTTGGT ACTITCGGTA GAAGCAACTG GTAAATTGTT AATTGGAACA 300 25 NTTGGTGGTG TAGATAACCA CGTATGGCCA AACCTAGAGC ATCTAGGCTC ACAATTACTA TCCTGACTTG ATAACAAGTG TTCTGATATT AACCTGAAAA TGGGAATAAT GCCAAATCTG 420 TGTAACTTAA CATCTATATA CACAGTGGGG AGAACTGAAG TTATTAAACC TGGAATCTCT 480 30 GTGATCAAGG CTAACAGTAG TTATCTAAGA AGCAAAGGAC CTACAATTCT TAGACTTGGA GTCATATTCT TTAAGGACGT GTTCTGAAAC TATATCAAGC ATCTGGTTTC CACGTATTTC 600 35 TCCCTCAGAA ATTATGAAGT ACAAGTAAAA ATGAAGGTAC AGGGTAAGAC ACATGCTGCT TTCTTGCTCT TGAGTGGAGA CAGTTTTCCA GCCATCTTAA CCCCTTWACA CAAAACAATT 720 TGTGTTTTAT AGCAAATAAG TGACTCAACA TAATTTCAAT ATGATGTTTA TCCACCAGTA 780 40 CTTTCCTTTC AGCTTCTAGT CCCATAARTG GTTTGTGAAG TCATCGGTTA CATTAGCCAA 840 GATAGGCCTA GACTTGAAGT CTAGAATGTT TTTCCCACTA TATGCCAAAG TAGAATGTGG 900 45 GTATCTCAGG GTCATTTTTG TTGTTCAATT TCCCACCTGT ACAGTTGTTA TGATTCACTT 960 1020 1047 CGAGGGGGG CCCGGTACCC AAATCGC 50

(2) INFORMATION FOR SEQ ID NO: 24:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 990 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

5	TTGGAAAGGG T	CTAGCTCTT	TCTCATTCAC	CAACTATATT	AGAAGCACTT	GAGGGAAATT	60
	TACCACTCCA A	ATCCAAAGC	AATGAACAGT	CTTTTCTGGA	TGATTTTATT	GCCTGTGTCC	120
10	CAGGATCAAG T	CGTGGAAGG	CTTGCAAGGT	GGCTTCAGCC	AGATTCATAT	GCGGATCCTC	- 180
10	AGAAAACATC T	TTGATCCTG	GAATAAGGAT	GATATTCGTT	GTGGTTGGCC	TACCACCATA	240
	ACTGTTCAAA C	CAAAAGACCA	CTATCCCCAT	GTGGTACATG	TTCCCAATAT	GAAGGTAATT	300
15	ATAACTGGAT 1	TAAATTAGCA	GACATCTATA	TACTGGCTGC	AATGACTGAT	AAAATTTTAG	360
	AAATGCCAAG 1	rcctgagrgt	CCATTTGTTC	TACCCTCTTT	ATATAAAGGG	TGATGCTGAA	420
20	AGITTGTTIA I	AATGACTTGT	TTATATTAAT	TAGTCCCCAA	GTGTCCAAGT	TACACCTGTT	480
20	TTTTTTGTGA (STTTGTTCTT	TACATTTTGC	TACCTGTTAC	GGGGACTCAA	AGGAGGGATA	540
	AGAAAGTATC (CATCTAAAGA	GTGCTAGACA	CATACAGTGA	AGCCCCTCAA	TATGTATTGA	606
25	TTGAATAAAT (GCATGAAAGA	ATACATTTT	AAATTTTGTG	TATAGTTTTG	AAAGACTCAA	66
	GTACGITCTG	IGTTTGGTAT	TACTGAAACC	ACATTTTAAA	AATAACACTC	ATTAAGTTAG	72
30	AAATATATGA (GTTTAGATTG	TAAAAGAATG	AGGAATTGAA	ATAGTTGTAT	ACCATATTGA	78
30	TGAATATAGA	GTTTTTAGGA	TACCTCTTAC	CTGAAATATT	AATAATAATG	TTTNCAGAGC	84
	ATATTATACA	TAATTATTT G	TGATTTAATC	TGTTAATATG	AATATCTCAT	TTAAAACTTT	90
35	TATTTCTGAA .	ТАТАТТААА	TGAATAAAAT	TTTATATAGG	CAGTCCCCAG	CCCTTTCCTC	96
	CTTCAAAGTT	GTCTTATAGA	GTGATTGGTT	•			99

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(2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1208 base pairs(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

60	CCGGGTCGAC	GTCCGGAATT	GCAGGTACCG	CCTCCTCCCT	TATAGGGAAA	TAATCGCTAC
120	GCTCCGGG	GCCTCCGCC	accesses	GCGCCTCCGG	GAGCGAAATG	CCACGCGTCC
180	CAGCAGTGCA	CGGCAGCTAC	CCTTCTACAT	GTAAAGAACG	GCTGTTCGAC	AGGTAGACGA
240	GGGACGTCTT	GACGTGGAGA	CCCAGAGAGA	AGCTRTCAAG	GCASGGGTGA	TAAACGAGGC
300	AGATCAAGCC	GTCCTGGATG	GTTCCGTGTG	CGCAGAGGAA	GCGTACCTGG	CCTGTATAGA

CTCCTCGCCC CCTGAGCTCC AGGCCGTGCG CATGTTTGCT GACTACCTCG CCCACGAGAG

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_	TCGGAGGGAC AGCATCGTGG CCGAGCTGGA CCGAGAGATG AGCAGGAGCK TCGACGTGAC	420
5	CAACACCACC TICCTGCTCA TGGCCGCCTC CATCTATCTC CACGACCAGA ACCCGGATGC	480
	CGCCCTGCGT GCGCTGCACC AGGGGGACAG CCTGGAGTGC ACAGCCATGA CAGTGCAGAT	540
10	CCTGCTGAAG CTGGACCGCC TGGACCTCGC CCGGAAGGAG CTGAAGAGAA TGCAGGACCT	600
	GGACGAGGAT GCCACCCTCA CCCAGCTCGC CACTGCCTGG GTCAGCCTGG CCACGGGTGG	660
	TGAGAAGCTG CAGGATGCCT ACTACATCTT CCAGGAGATG GCTGACAAGT GCTCGCCCAC	720
15	CCTGCTGCTG CTCAATGGGC AGGCGGCCTG CCACATGGCC CAGGGCCGCT GGGAGGCCGC	780
	TGAGGGCCTG CTGCAGGAGG CGCTAGACAA GGATAGTGGC TACCCRGAGA CGCTGGTCAA	840
20	CCTCATCGTC CTGTCCCAGC ACCTKGGCAA GCCCCCTGAG GTGACAAACC GATACCTGTC	900
	CCAGCTGAAG GATGCCCACA GGTCCCATCC CTTCATCAAG GAGTACCAGG CCAAGGAGAA	960
25	CGACTTIGAC AGGCTGGTGC TACAGTACGC TCCCAGCGCT GAGGCTGGCC CAGAGCTGTC	1020
25	AGGACCATGA AGCCAGGACA GAGGCCAGGA GCCAGCCCTG CAGCCCTCCC CACCCGGCAT	1080
	CCACCTGCAT CCCTCTGGGG CAGGAGCCCA CCCCCAGCAC CCCCATCTGT TAATAAATAT	1140
30	CTCAACTCCA RGGTGTTCCA CCTGAAAAAA AAAAAAAAA AAAAAAAAA AAAAAAAAA	1200
	аааааааа -	1208
35		
33	(a) THOMOTON TON GEO ID NO. 26.	
	(2) INFORMATION FOR SEQ ID NO: 26:	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1922 base pairs	
	(B) TYPE: nucleic acid	
٠.	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:	
	GTGCTGCGCT ACTGAGCAGC GCCATGGAGG ACTCTGAAGC ACTGGGCTTC GAACACATGG	60
50	GCCTCGATCC CCGCCTCCTT CAGGCTGTCA CCGATCTGGG CTGGTCGCGA CCTACGCTGA	120
50	TCCAGGAGAA GGCCATCCCA CTGGCCCTAG AAGGGAAGGA CCTCCTGGCT CGGGCCCGCA	180
	CGGGCTCCGG GAAGACGGCC GCTTATGCTA TTCCGATGCT GCAGCTGTTG CTCCATAGGA	240
55	AGGCGACAGG TCCGGTGGTA GAACAGGCAG TGAGAGGCCT TGTTCTTGTT CCTACCAAGG	300
	ACCURACION CONNECTOR TOCATGATTO AGCAGOTGGO TACCTACTGT GOTGGGGATG	360

TCCGAGTGCC CAATGTCTCA GCTGCTGAAG ACTCAGTCTC TCAGAGAGCT GTGCTGATGG

•	AGAAGCCAGA TGTGGTAGTA GGGACCCCAT CTCGCATATT AAGCCACTTG CAGCAAGACA	480
	GCCTGAAACT TCGTGACTCC CTGGAGCTTT TGGTGGTGGA CGAAGCTGAC CTTCTTTTTT	540
5	CCTTTGCCTT TGAAGAAGAG CTCAAGAGTC TCCTCTGTCA CTTGCCCCGG ATTTACCAGG	600
	CITTITCICAT GTCAGCTACT TTTAACGAGG ACGTACAAGC ACTCAAGGAG CTGATATTAC	660
10	ATAACCCGGT TACCCTTAAG TTACAGGAGT CCCAGCTGCC TGGGCCAGAC CAGTTACAGC	- 720
10	AGTITICAGGI GGICIGIGAG ACTGAGGAAG ACAAATICCI CCTGCTGTAT GCCCTGCTCA	780
	AGCTGTCATT GATTCGGGGC AAGTCTCTGC TCTTTGTCAA CACTCTAGAA CGGAGTTACC	840
15	GGCTACGCCT GTTCTTGGAA CAGTTCAGCA TCCCCACCTG TGTGCTCAAT GGAGAGCTTC	900
	CACTGCGCTC CAGGTGCCAC ATCATCTCAC AGTTCAACCA AGGCTTCTAC GACTGTGTCA	960
20	TAGCAACTGA TGCTGAAGTC CTGGGGGCCC CAGTCAAGGG CAAGCGTCGG GGCCGAGGGC	1020
20	CNAAAGGGGA CAAGGCCTCT GATCCGGAAG CAGGTGTGGC CCGGGGCATA GACTTCCACC	1080
	ATGIGTCTGC TGTGCTCAAC TTTGATCTTC CCCCAACCCC TGAGGCCTAC ATCCATCGAG	1140
25	CTGGCAGGAC AGCACGCGCT AACAACCCAG GCATAGTCTT AACCTTTGTG CTTCCCACGG	1200
	AGCAGTTCCA CTTAGGCAAG ATTGAGGAGC TTCTCAGTGG AGAGAACAGG GGCCCCATTC	1260
30	TGCTCCCCTA CCAGTTCCGG ATGGAGGAGA TCGAGGGCTT CCGCTATCGC TGCAGGGATG	1320
30	CCATGCGCTC AGTGACTAAG CAGGCCATTC GGGAGGCAAG ATTGAAGGAG ATCAAGGAAG	1380
	AGCTTCTGCA TTCTGAGAAG CTTAAGACAT ACTTTGAAGA CAACCCTAGG GACCTCCAGC	1440
35	TGCTGCGGCA TGACCTACCT TTGCACCCCG CAGTGGTGAA GCCCCACCTG GGCCATGTTC	1500
	CTGACTACCT GGTTCCTCCT GCTCTCCGTG GCCTGGTRCG CCCTCACAAG AAGCGGAAGA	1560
40	AGCTGTCTTC CTCTTGTAGG AAGGCCAAGA GAGCAAAGTC CCAGAACCCA CTGCGCAGCT	1620
	TCAAGCACAA AGGAAAGAAA TTCAGACCCA CAGCCAAGCC CTCCTGAGGT TGTTGGGCCT	1680
	CTCTGGAGCT GAGCACATTG TGGAGCACAG GCTTACACCC TTCGTGGACA GGCGAGGCTC	1740
45	TGGTGCTTAC TGCACAGCCT GAACAGACAG TTCTGGGGCC GGCAGTGCTG GGCCCTTTAG	1800
	CTCCTTGGCA CTTCCAAGCT GGCATCTTGC CCCTTGACAA CAGAATAAAA ATTTTAGCTG	1860
50	CCCCAAAAAA AAAAAAAAAA AAAAAAACTC GAGGGGGGC CCGTACCCAA TTCGCCCTAT	1920
50	AA	1922

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1951 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

5 TOSTCCCCAG AGCGGGCTGA GCCCCAGGCG SAGGGTGGCG GGGGAGCCTG GGGGAGCCGC 60 COCCACCTCC ACGGGCCTCT CTGAGCTCGG ACACCAGCGC CCTGTCCTAT GACTCTGTCA 120 AGTACACGCT GGTGGTAGAT GAGCATGCAC AGCTGGAGCT GGTGAGCCTG CGCCGTGCTT 180 10 COGAGACTAC AGTGACGAGA GTGACTCTCC CACCGTCTAT GACAACTGTG CCTCCGTCTC 240 CTCGCCCTAT GAGTCGCCCA TCGGAGAGGA ATATGAGGAG GCCCCGCGGC CCCAGCCCCC 300 15 TOCCTGCCTC TCCGAGGAAC TCCACGCCTG ATGAACCCGA CGTCCATTTC TCCAAGAAAT 360 TCCTGAACGT YTTCATGAGT GGCCGCTCCC GCTCCTCCAG TGCTGAGTCC TTCGGGCTGT 420 TCTCCTGCAT CATCAACGGG GAGGAGCAGG AGCAGACCCA CCGGGCCATA TTCAGGTTTG 480 20 540 TGCCTCGACA CGAAGACGAA CTTGAGCTGG AAGTGGATGA CCCTCTGCTA GTGGAGCTCC AGGCTGAAGA CTACTGGTAC GAGGCCTACA ACATGCGCAC TGGTGCCCGG GGTGTCTTTC 600 25 660 CTGCCTATTA CGCCATCGAG GTCACCAAGG AGCCCGAGCA CATGGCAGCC CTGGCCAAAA ACAGTGACTG GGTGGACCAG TTCCGGGTGA AGTTCCTGGG CTCAGTCCAG GTTCCCTATC 720 ACAAGGGCAA TGACGTCCTC TGTGCTGCTA TGCAAAAGAT TGCCACCACC CGCCGGCTCA 30 CCGTGCACTT TAACCCGCCC TCCAGCTGTG TCCTGGAGAT CAGCGTGCGG GGTGTGAAGA 840 TAGGCGTCAA GGCCGATGAC TCCCAGGAGG CCAAGGGGAA TAAATGTAGC CACTTTTTCC 900 35 AGTTAAAAAA CATCTCTTTC TGCGGATATC ATCCAAAGAA CAACAAGTAC TTTGGGTTCA 960 TCACCAAGCA CCCCGCCGAC CACCGGTTTG CCTGCCACGT CTTTGTGTCT GAAGACTCCA 1020 1080 CCAAAGCCCT GGCAGAGTCC GTGGGGAGAG CATTCCAGCA GTTCTACAAG CAGTTTGTGG 40 AGTACACCTG CCCCACAGAA GATATCTACC TGGAGTAGCT GTGCAGCCCC GCCCTCTGCG 1140 TCCCCCAGCC CTCAGGCCAG TGCCAGGACA GCTGGCTGCT GACAGGATGT GGCACTGCTT 1200 45 GAGGAGGGG ACCTGCCACC GCCAGAGGAC AAGGAAGTGG GGCGCTGGCC CAGGGTAGGG 1260 1320 GAGGGTGGGG CAATGGGGAG AGGCAAATGC AGTTTATTGT AATATATGGG ATTAGATTCA TCTATGGAGG GCAGAGTGGG CTGCCTGGGG ATTGGGAGGG ACAGGGCTTG GGGAGCAGGT 50 1380 CTCTGGCAGA GAAGGATGTC CGTTCCAGGA GCACACGGCC CTGCCCCATC CTGGGCCTTA 1440 CCTCCCCTGC CAGGGCTCGG GCGCTGTGGC TCCTGCCTTG ATGAAGCCCG TGTCCTGCCT 1500 . 55 1560 TGATGAAGCC TGTGCCACCT GCAAGTGCCC GCCCTGCCCC TGCCCCAACC CCCACCGAAG AGCCCTGAGC TCAGGCTGAG CCCAGCCACC TCCCAAGGAC TTTCCAGTGA GGAAATGGCA 1620 1680 ACACGIGGAG GIGAAGICCC TGITCTCAGC TCCGTCATCT GCGGGGCTIC TGGGIGGCTC 60

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-	CTGCCACTGA	CCTCACCGGC	ATGCTGGCCT	GTGGCAGGCC	TAGGACCTCA	GGCGGGGAGG	1740
_	AGGAGCTGCC	GCAAGGCCCT	GTCCCAGCAG	AAGAGGGAGG	CTTCCTGACT	GACACAGGCC	1800
5	AGCCCCATCT	TOGTCCTGTC	ACCCTGGCCC	CAACTATTAA	AGTGCCATTT	CCTGTCAAAA	1860
	ааааааааа	AAAATCGGGG	GGGGCCCGGA	ANCCAATTTC	CCCCAAAAAG	GGGGGTTATA	1920
10	AAAATTCCCN	GCCNGTGTTT	TTAAAAATTC	G			1951

(2) INFORMATION FOR SEQ ID NO: 28: 15

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3989 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

25	GGCACAGGCC GCAGGGNACC TATGGGCGCA TATAGGTTGT AATGAAACTG TAGTCTCAGT	60
	TEGAAGCCTA GACATGAAAT GEGTCAGTGA GCAAGGCTCT ATTCCTAGTC TCCAGCCATG	120
30	CCTGTGGAAC CTGARCCCRC TCTCAGCACA TTGGACCCAG GCAGATGYAA AAAATTCACA	180
	GAACTATGAT TIGGACTCAA GGGITTGIAG ATTICCTCCT TCATTCTAAT TICAGIGICT	240
	AAAATTCTTG CATCCRTGAA CGAGCTGGGC ATTTGATGAG ACAGGGCYGA ATACTGCAGT	, 300
35	TITICCTCCTA GAAATCATCT GGGGCATTTT CTTTGAACTG ATGGGAACAA TAAGGCATAA	360
	CTGTTTGCAC AAACTTGGGA TAARTGATTT TGGGATAACG ATCTACCAGA ATGGGGATAT	420
	TTCACCCTTG GTTCTGAGAT GCAAACCAAA GAATATCATG ACCAGCTTTC AGGCCTCCTG	480
40	AAGTATATCT CTCACATTGT CCTGTTCTCA TGCTGAGGAG CCTGAGATCC CTGTGTGGGG	540
	ATTAGACAGT GGACTGTTAT GGGTGTAGGT GAATTGGCTT ATTTTGTCTG TCCCTGTCTG	600
45	AATGTATTGC AGGAAYTAAA AAGGACCAAG AAGAGGAAGA AGACCAAGGC CCACCATGCC	660
	CCAGGCTCAG CAGGGAGCTG CTGGAGGTAG TAGAGCCTGA AGTCTTGCAG GACTCACTGG	720
50	ATAGATGITA TICAACICCI TCCAGITGIC TIGAACAGCC TGACTCCTGC CAGCCCTATG	780
	GAAGTICCIT TTATGCATTG GAGGAAAAAC ATGTTGGCTT TTCTCTTGAC GTGGGAGAAA	840
	TTGAAAAGAA GGGGAAGGGG AAGAAAAGAA GGGGAAGAA	900
	GGGGAAGAAA AGAAGGGGAA GAAGATCAAA ACCCACCATG CCCCAGGCTC AGCAGGGAGC	960
<i>)</i>	TECTOGATGA GAAAGRECCT GAAGTCTTGC AGGACTCACT GGATAGATGT TATTCAACTC	1020
	CTTCAGTTGT GTTGAACTGT GTGACTCATG CCAGCCCTAC AGAAGTGCCT TTTATGTATT	1080
60	CTICAGTIGT GITGAACIGT GIGACTCATG CCAGCCCTAC ABARDIOCCT	

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•	GGAGCAACAG CATGTTGGCT TGGCTGTTGA CATGGATGAA ATTGAAAAGT ACCAAGAAGT	1140
	GGAAGAAGAC CAAGACCCAT CATGCCCCAG GCTCAGCAGG GAGCTGCTGG ATGAGAAAGA	1200
5	GCCTGAAGTC TTGCAGGACT CACTGGATAG ATGTTATTCG ACTCCTTCAG GTTATCTTGA	1260
	ACTOCCTGAC TTAGGCCAGC CCTACAGCAG TGCKGTTTAC TCATTGGAGG AMCAKTACCT	1320
10 15	TGGCTTKKCT CTTGACGTGG ASAAATTGAA AAGAAGGGGA AGGGGAARAA AAGAAGGGGA .	1380
	AGAAGATCAA AGAAGGAAAG AAGAAGGGGA AGAAAAGAAG GGGAAGAA	1440
	CCATGCCCCA GGCTCAGCAG GGAGCTGCTG GATGAGAAAG GGCCTGAAGT CTTGCAGGAC	1500
	TCACTGGATA GATGITATIC AACTCCTTCA GGTTGTCTTG AACTGACTGA CTCATGCCAG	1560
	CCCTACAGAA GTGCCTTTTA YRTATTGGAG CAACAGYGTG TTGGCTTGGC TGTTGACATG	1620
20	GATGAAATTG AAAAGTACCA AGAAGTGGAA GAAGACCAAG ACCCATCATG CCCCAGGCTC	1680
	AGCAGGGAGC TGCTGGATGA GAAAGAGCCT GAAGTCTTGC AGGACTCACT GGATAGATGT	1740
25	TATTCGACTC CTTCAGGTTA TCTTGAACTG CCTGACTTAG GCCAGCCCTA CAGCAGTGCT	1800
	GTTTACTCAT TGGAGGAACA GTACCTTGGC TTGGCTCTTG ACGTGGACAG AATTAAAAAG	1860
	GACCAAGAAG AGGAAGAAGA CCAAGGCCCA CCATGCCCCA GGCTCAGCAG GGAGCTGCTG	1920
30	GAGGTAGTAG AGCCTGAAGT CTTGCAGGAC TCACTGGATA GATGTTATTC AACTCCTTCC	1980
	AGITGICITG AACAGCCIGA CICCIGCCAG CCCIATGGAA GITCCITITA IGCATIGGAG	2040
35	GAAAAACATG TTGGCTTTTC TCTTGACGTG GGAGAAATTG AAAAGAAGGG GAAGGGGAAG	2100
	AAAAGAAGGG GAAGAAGATC AAMGAAGRAA AGAAGAAGGG GAAGAAAAGA AGGGGAAGAA	2160
	GATCAAAACC CACCATGCCC CAGGCTCAAC GGCGTGCTGA TGGAAGTGGA AGAGCSTGAA	2220
40	GTCTTACAGG ACTCACTGGA TAGATGTTAT TCGACTCCGT CAATGTACTT TGAACTACCT	2280
	GACTCATTCC AGCACTACAG AAGTGTGTTT TACTCATTTG AGGAACAGCA CATCAGCTTC	2340
45	GCCCTTTACG TGGACAATAG GTTTTTTACT TTGACGGTGA CAAGTCTCCA CCTGGTGTTC	2400
	CAGATGGGAG TCATATTCCC ACAATAAGCA GCCCTTASTA AKCCGAGAGA TGTCATTCCT	2460
	GCAGGCAGGA CCIATAGGCA MGTGAAGATT TGAATGAAAG TACAGTTCCA TTTGGAAGCC	2520
50	CAGACATAGG ATGGGTCAGT GGGCATGGCT CTATTCCTAT TCTCAAACCA TGCCAGTGGC	2580
	AACCTGTGCT CAGTCTGAAG ACAATGGACC CACGTTAGGT GTGACACGTT CACATAACTG	2640
	TGCAGCACAT GCCGGGAGTG ATCAGTCRGA CATTTTAATT TGAACCACGT ATCTCTGGGT	2700
55	AGCTACAAAA TTCCTCAGGG ATTTCATTTT GCAGGCATGT CTCTGAGCTT CTATACCTGC	276
	TCAAGGTCAK TGTCATCTTT GTGTTTAGCT CATCCAAAGG TGTTACCCTG GTTTCAATGA	282
60	ACCTAACCTC ATTCTTTGTG TCTTCAGTGT TGGCTTGTTT TAGCTGATCC ATCTGTAACA	288
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	CAGGAGGGAT CCTTGGCTGA GGATTGTATT TCAGAACCAC CAACTGCTCT TGACAATTGT	2940
	TAACCCCCTA GRCTCCTTTG GTTAGAGAAG CCACAGTCCT TCAGCCTCCA ATTGGTGTCA	3000
5	GTACTTAGGA AGACCACAGC TAGATGGACA AACAGCATTG GGAGGCCTTA GCCCTGCTCC	3060
	TCTCRATTCC ATCCTGTAGA GAACAGGAGT CAGGAGCCGC TGGCAGGAGA CAGCATGTCA	3120
10	CCCAGGACTC TGCCGGTGCA GAATATGAAC AAYGCCATGT TCTTGCAGAA AACGCTTAGC	3180
10	CTGAGTTTCA TAGGAGGTAA TCACCAGACA ACTGCAGAAT GTRGARCACT GAGCAGGACA	3240
	GCTGACCTGT CTCCTTCACA TAGTCCATRT CACCACAAAT CACACAACAA AAAGGAGARG	3300
15	AGATATTITG GGITCAAAAA AAGTAAAAAG ATAATGTAGC TGCATTTCTT TAGITATTIT	3360
	GARCCCCAAA TATTTCCTCA TCTTTTTGTT GTTGTCATKG ATGGTGGTGA CATGGACTTG	3420
20	TTTATAGAGG ACAGGTCAGC TGTCTGGCTC AGTGATCTAC ATTCTGAAGT TGTCTGAAAA	3480
20	TGTCTTCATG ATTAAATTCA GCCTAAACGT TTTGCCGGGA ACACTGCAGA GACAATGCTG	3540
	TGAGTTTCCA ACCTYAGCCC ATCTGCGGGC AGAGAAGGTC TAGTTTGTCC ATCASCATTA	3600
25	TCATGATATC AGGACTGGTT ACTTGGTTAA GGAGGGGTCT AGGAGATCTG TCCCTTTTAG	3660
	AGACACCITA CTTATAATGA AGTATTTGGG AGGGTGGTTT TCAAAATTAG AAATGTCCTG	3720
30	TATTCCRATG ATCATCCTGT AAACATTTTA TCATTTATTA ATCATCCCTG CCTGTGTCTA	3780
50	TTATTATATT CATATCTCTA COCTGGAAAC TTTCTGCCTC AATGTTTACT GTGCCTTTGT	3840
	TTTTGCTAGT GTGTGTTGTT GAAAAAAAA ACATTCTCTG CCTGAGTTTT AATTTTTGTC	3900
35	САААСТТАТТ ТТААТСТАТА СААТТААААС СТТТТСССТА ТСАААААААА	3960
	AAAAAAAAA AAAAAGCGGA CGCGTGGGC	3989

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(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3735 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

60	GTCGGCGGGC	CSGCTGACGG	GCTTGGCCAG	TCCGCAGCAG	CTGGCTGGGC	CIGCIGITCG
120	CCCTCTCAAA	GGTAGTGCAN	ATTTTATTCT	GCAGCTGCAG	GAACAGGCAC	GGGTTTGTGT
180	AGAAAACTIG	ATTCCAAAAA	AGAAGTAGTA	CAGGGATTGA	ACTGATGTAA	GGTTGAAGGA
240	A CCACAGCTGT	AACAGGGATA	ATCCACAGTA	AGGCACTTGC	GCCGTTCTTC	GGATAAAGTA
300	AATCTCGTTC	TCATCTTTGG	TATGCCAGCA	ATCCTTACCT	TTTCAAGATG	GCCTTATGTG

	ATTITIACTG GCAAAGAAAT CCGGGGAGAA TGTGGCCAAG TTTATTATTA ATTCATACCC	360
5	CARATATITT CAGAAGGACA TAGCTGAACC TCATATACCG TGTTTAATGC CTGAGTACTT	420
	TGAACCTCAG ATCAAAGACA TAAGTGAAGC CGCCCTGAAG GAACGAATTG AGCTCAGAAA	480
	AGTCAAAGCC TCTGTGGACA TGTTTGATCA GCTTTTGCAA GCAGGAACCA CTGTGTCTCT	540
10	TGAAACAACA AATAGTCTCT TGGATTWIT GTGTTACTAT GGTGACCAGG AGCCCTCAAC	600
	TGATTACCAT TTTCAACAAA CTGGACAGTC AGAAGCATTG GAAGAGGAAA ATGATGAGAC	660
15	ATCTAGGAGG AAAGCTGGTC ATCAGTTTGG AGTTACATGG CGAGCAAAAA ACAACGCTGA	720
15	GAGAATCTTT TCTCTAATGC CAGAGAAAAA TGAACATTCC TATTGCACAA TGATCCGAGG	780
	AATGGTGAAG CACCGAGCTT ATGAGCAGGC ATTAAACTTG TACACTGAGT TACTAAACAA	840
20	CAGACTCCAT GCTGATGTAT ACACATTTAA TGCATTGATT GAAGCAACAG TATGTGCGAT	900
	AAATGAGAAA TTTGAGGAAA AATGGAGTAA AATACTGGAG CTGCTAAGAC ACATGGTTGC	960
25	ACAGAAGGTG AAACCAAATC TICAGACTIT TAATACCATT CIGAAATGTC TCCGAAGATT	1020
23	TCATGTGTTT GCAAGATCGC CAGCCTTACA GGTTTTACGT GAAATGAAAG CCATTGGAAT	1080
	AGAACCCTCG CTTGCAACAT ATCACCATAT TATTCGCCTG TTTGATCAAC CTGGAGACCC	1140
30	TITAAAGAGA TCATCCTTCA TCATTTATGA TATAATGAAT GAATTAATGG GAAAGAGATT	1200
	TICTCCAAAG GACCCGGATG ATGATAAGIT TITTCAGICA GCCATGAGCA TATGCTCATC	1260
35	TCTCAGAGAT CTAGAACTIG CCTACCAAGT ACATGGCCTT TTAAAAACCG GAGACAACTG	1320
33	GAAATTCATT GGACCTGATC AACATCGTAA TTTCTATTAT TCCAAGTTCT TCGATTTGAT	1380
	TTGTCTAATG GAACAAATTG ATGTTACCTT GAAGTGGTAT GAGGACCTGA TACCTTCAGC	1440
40	CTACTITCCC CACTCCCAAA CAATGATACA TCTTCTCCAA GCATTGGATG TGGCCAATCG	1500
	GCTAGAAGTG ATTCCTAAAA TTTGGAAAGA TAGTAAAGAA TATGGTCATA CTTTCCGCAG	1560
45	TGACCTGAGA GAAGAGATCC TGATGCTCAT GGCAAGGGAC AAGCACCCAC CAGAGCTTCA	1620
43	GGTGGCATTT GCTGACTGTG CTGCTGATAT CAAATCTGCG TATGAAAGCC AACCCATCAG	1680
	ACAGACTECT CAGGATTEGC CAGCCACCTC TCTCAACTET ATAGCTATCC TCTTTTTAAG	1740
50	GCCTGGGAGA ACTCAGGAAG CCTGGAAAAT GTTGGGGCTT TTCAGGAAGC ATAATAAGAT	1800
	TCCTAGAAGT GAGTTGCTGA ATGAGCTTAT GGACAGTGCA AAAGTGTCTA ACAGCCCTTC	1860
55	CCAGGCCATT GAAGTAGTAG AGCTGGCAAG TGCCTTCAGC TTACCTATTT GTGAGGGCCT	1920
55	CACCCAGAGA GTAATGAGTG ATTTTGCAAT CAACCAGGAA CAAAAGGAAG CCCTAAGTAA	1980
	TCTAACTGCA TTGACCAGTG ACAGTGATAC TGACAGCAGC AGTGACAGCG ACAGTGACAC	2040
60	CACTGARGE ANATCANET GGAGATTCAG GAGCAGCAAT GGTCTCACCA TAGCTGCTGG	2100

•	AATCACACCT GAGAACTGAG ATATACCAAT ATTTAACATT GTTACAAAGA AGAAAAGATA	2160
5	CAGATTTGGT GAATTTGTTA CTGTGAGGTA CAGTCAGTAC ACAGCTGACT TATGTAGATT	2220
	TAAGCTGCTA ATATGCTACT TAACCATCTA TTAATGCACC ATTAAAGGCT TAGCATTTAA	2280
	GTAGCAACAT TGCGGTTTTC AGACACATGG TGAGGTCCAT GGCTCTTGTC ATCAGGATAA	2340
10	GCCTGCACAC CTAGAGTGTC GGTGAGCTGA CCTCACGATG CTGTCCTCGT GCGATTGCCC	2400
	TCTCCTGCTG CTGGACTTCT GCCTTTGTTG GCCTGATGTG CTGCTGTGAT GCTGGTCCTT	2460
1.5	CATCITAGGT GITCATGCAG TICTAACACA GITGGGGITG GGTCAATAGT TICCCAATIT	2520
15	CAGGATATTT CGATGTCAGA AATAACGCAT CTTAGGAATG ACTAAACAAG ATAATGGCAG	2580
	TTTAGGCTGC ACAACTGGTA AAATGACTGT AGATAAATGT TGTAATTAGT GTACACGTTT	2640
20	GTATTTTGT TAATATAGCC GCTGCCATAG TTTTCTAACT TGAACAGCCA TGAATGTTTC	2700
	ATGTCTCCCT TTTTTTTTTG TCTATAGCTG TTACCTATTT TAGTGGTTGA AATGAGAGCT	2760
25	AGTGATGACA GAAGGATGTG GAATGTCTTC TTGACATCAT TGTGTATTGC TGGTAATCAA	2820
23	GTTGGTAACG ACTACTTCTA GCAGCTCTTA CCACTATGAC TTAAGTGGTC CTGGAAGGCA	2880
	GTAAGTGGAG GTTTGCAGCA TTCCTGCCTT CATGAGGGCT TCTACCACTG ACCACTTTGC	2940
30	ACGTACCTGG CTCCCAGATT TACTTAGGTA CCCCACGAGT CGTCCACATA AGCAGCTTCA	3000
	TOTTTACCTT GCCAGAGTTG ACAATTATGG GATACTCTAG TCTACTTATA CTTGTGTTCC	3060
35	CATCTGTCTG CCATCCTCTG AAGGCCAGGA CCCAGTCATA CATCCTTAGA AACCAAAGTA	3120
,,,	TOGITTITIGI TITCTCTTGG AATGTCAGGT CTTAAGGCAT TTAATTGAGG GACAAAAAAA	3180
	AAAAAAAGCC GATATAGTAG CTAGCTACTT AAGCATCCAT GGGTATTGCT CCATATCAAA	3240
40	GCAGATTTGC AGGACAGAAA GAGTAAATTA GCCTTCAGTC TTGGTTTACA GCTTCCAAGG	3300
	AGAGCCTTGG CCACCTGAAA TGTTAACTCG GTCCCTTCCT GTCTCTAGTT CATCAGCACC	3360
45	TGCAGATGCC TGACTCTTGT TAGCCTTACT ATTCAATACA GTCCTTAGAT TCACGGTATG	3420
73	CCTCTTCCTA TCCAGGCACC TATTCTGAAT CACCATGTTG CTCTGCAGCT AGAGTTGATA	3480
	GGAGAAAATC CATTTGGGTA GATGGCCTAT GAATTTGTAG TAGACTTTCA AAATGAGTGA	3540
50	TTTGTTAGCT TGGTACTTTT AAGTTTGTGG TACAGATCCT CCAAACCCAT ACTCTGAGCA	3600
	ATTAACTGCC TTGAACATAG AGAAAATTAA GGCCTCACAG GATGAGTCTC CATTCTCTGT	3660
55	AAATGCTTAT TTTATCATAG TCTTTAGCCN CTACTATGAG TAAAATGTTC TCTTCNGCCG	3720
55	GGTGTGGTGA CTCAC	3735

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(2) INFORMATION FOR SEQ ID NO: 30:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1667 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

10 TAGTAATICA TITAACICCT CTTACATGAG TAGCGACAAT GAGTCAGATA TCGAAGATGA 60 AGACTTAAAG TTAGAGCTGC GACGACTACG AGATAAACAT CTCAAAGAGA TTCAGGACCT 120 GCAGAGTCGC CAGAAGCATG AAATTGAATC TTTGTATACC AAACTGGGCA AGGTGCCCCC 180 15 TGCTGTTATT ATTCCCCCAG CTGCTCCCCT TTCAGGGAGA AGACGACGAC CCACTAAAAG 240 CARAGGCAGC ARATCTAGTC GAAGCAGTTC CTTGGGGART ARAAGCCCCC AGCTTTCAGG 300 20 TAACCTGTCT GGTCAGAGTG CAGCTTCAGT CTTGCACCCC CAGCAGACCC TCCACCCTCC 360 TGGCAACATC CCAGAGTCCG GGCAGAATCA GCTGTTACAG CCCCTTAAGC CATCTCCCTC 420 480 CAGTGACAAC CTCTATTCAG CCTTCACCAG TGATGGTGCC ATTTCAGTAC CAAGCCTTTC 25 TOCTCCAGGT CAAGGAACCA GCAGCACAAA CACTGTTGGG GCAACAGTGA ACAGCCAAGC 540 CGCCCAAGCT CAGCCTCCTG CCATGACGTC CAGCAGGAAG GGCACATTCA CAGATGACTT 600 30 GCACAAGTTG GTAGACAATT GGGCCCGAGA TGCCATGAAT CTCTCAGGCA GGAGAGGAAG 660 CAAAGGCAC ATGAATTATG AGGGCCCTGG AATGGCAAGG AAGTTCTCTG CACCTGGGCA 720 ACTOTOCATC TECATGACET CGAACETGGG TGGCTCTGCC CCCATCTCTG CAGCATCAGC 780 35 TACCTOTOTA GGTCACTTCA CCAAGTCTAT GTGCCCCCCA CAGCAGTATG GCTTTCCAGC 840 TACCCCATTT GGCGCTCAAT GGAGTGGGAC GGGTGGCCCA GCACCACAGC CACTTGGCCA 40 GTTCCAACCT GTGGGAACTG CCTCCTTGCA GAATTTCAAC ATCAGCAATT TGCAGAAATC 960 CATCAGCAAC CCCCCAGGCT CCAACCTGCG GACCACTTAG ACCTAGAGAC ATTAACTGAA 1020 TAGATCTGGG GGCAGGAGAT GGAATGCTGA GGGGTGGGT GGGGGTGGGA AGTAGCCTAT 1080 45 ATACTAACTA CTAGTGCTGC ATTTAACTGG TTATTTCTTG CCAGAGGGGA ATGTTTTTAA 1140 TACTGCATTG ACCCCTCAGA ATGGAGAGTC TCCCCCGCTC CAGTTATTGG AATGGGAGAG 1200 50 GAAGGAAAGA ACAGCTTTTT TGTCAAGGGG CAGCTTCAGA CCATGCTTTC CTGTTTATCT 1260 ATACTCAGTA ATGAGGATGA GGGCTAGGAA AGTCTTGTTC ATAAGGAAGC TGGAGAACTC 1320 1380 AATGTAAAAT CAAACCCATC TGTAATTTCG AGTGGGTGGA GCTCTTGCTT TTGGTACATG 55 CCCTGAATCC CTCACTCCCT CAAGAATCCG AACCACAGGA CAAAAACCAC CTACTGGGCT 1440 CTCTCCTACC CTGCCCTCCT CCCTTTTTT TACCCCTCTC TTTTTTATTT TTTCTTTGCT 1500 WO 98/54963

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CTTTAGAACC	CAGTGAAAAA	TACCAGGGTA	CTGGGGTGCA	ACTOTTTCTT	ATGATAGGTC	1560
ATTAGTGCTT	TAAGCAAAAG	ATATTAGCAG	CTTTGACTGC	AGCATTAGCA	ATTAGGRAAA	1620
************************	AAAACTYCAG	cccccccc	GTTACCCAAT	TCGCCCT		1667

10 (2) INFORMATION FOR SEQ ID NO: 31:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1408 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

20	ATTACACACC TGAGCACTGT GCCTGGCAAG ACCTGTCTTA ATAGATTAGA GAACCACTGA	60
	TAGATGGTCA GCTTTCTGTA GCAGTGAGAA CCCTACATTT CAAATGTGGA TAGCACCTTT	120
	GCGGGGAAAC ATCACTTGGC ACATCTGCAT TCTTTTTTGA CACAGGGTCT CACTCTGTTG	180
25	CCCAGGCTAG AGTGCATGGC ACGATCTTAG CTCACTGCAA CCTCCACCTC CCAAGTTCAA	240
	GCGATTCTTC TGCCTCAGCC TCCTGAGCAG CTGGGATCAC AGACATGCGC TACCATGCCC	300
30	AGCTAATITT TIGTATTITT TGTKTGTTTG TTTTTGTTTK TAAGTAGAGA CGGGCTTTCA	360
	CCACGTTGGS CAGGCAGGTC TCGAACTCCT GAMCTCAGGT GATCCACCCA CATCTGCGTT	420
	CCAATATCTT TCTCAACATA ATGATAGCCG TAATTAATAT TTTCCAGTAC ATTTTTATGC	480
35	CTTTACACAC GAGAGTOGTA GACAGACACA AACCCAGATC TGTCTGACTC CAAAGCCCGT	540
	TIGTCATCAT TOCTITTACG GTATCCTATA GTGGTATCCT TTACAGAAAG ACAGCTTTTA	600
40	CCCAACAAG ACTTAACTTC CCAGGATGCC AGAAGGACAA AGCGGGATTG CTTTTAAGRA	660
	GRAAGTTATC AAGAMCTTAT TITATAAATG AGATTAGATA GGGAAAGGCA ATTTATCTTT	720
	ATTAAAAACT GAAAAGGCCA GCATAGGGAA GGAGGTCCTT CGGTGGTCTT TTTCAGGGAA	780
45	ATACTTCAGT TGCTTTTATT AGAAACAGAT AGTACCTAAG GTTTTGAGGT AGGWACAGCT	840
	TAAGGCATGC TAATGKTCAT GGGTCCTTCC ATAGTCATTT TKGTATTTTG GTTWACATTT	900
50	GASCANTAGG CAGCCCTTCA CTGCTGCTGG AYTCATTCCT GCCAYTATTA CAGGTGACAG	960
	AGGAGACAGG AGGIATGICT TITCTATITT TAWACATGCT TTATATITAA CACAAGCTCT	1020
	TGGGTATCTT AGATAAACAG AAGTTGCCTA GCACTCCTTT TAGTGCATTG AACCCTTTAA	1080
55	CATTTAAGCA AAATAATAAA CAGTCTTTTG AGGTTCCTTA ACAATGAAAC GTGTTCGAGT	1140
	GGCAGCAGCG GAATCCATGC YTCTTCTCCT GGAGTGTGCA AKAGTCCGTG GTCCTGAGTA	1200
60	TCTCACACAG ATGTGGCATT TTATGTGTGA TGCTCTAATT AAGGCCATTG GTACAGAACC	1260

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AGATTCAGAC GTCCTCTCAG AAATAATGCA TTCTTTTGCA AAGGTGAATA TTTTTCTCTT	1320
AAAAAATATG TATAAGGTGG TATGTTCATT TATTAGTCTT GCTAAAAAAA AAAAAAAAAA	1380
ACTINGAGGG GGGGNCCGGT ACCCAATT	1408

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(2) INFORMATION FOR SEQ ID NO: 32:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2031 base pairs

15 (3) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:

20 AGGATATGCA TGATTCTTAA CCAGGCTATA TGTTAAAAAA AAATTGGAAA ATGCAATACA 60 TTTTTTATTA TACAAACTAC AGAATGAGTA TGCAAGTTTT ATTTATCAAA ATGTAATGGA 120 TITITAAAGG CIGAGAAATT TECCITATAC CIACCITITC AGITATITTA ATTATACCAA 180 25 ATTATCAACT AGAATAGCTT CATCCATATG AAATATAAAA TGAAGAGACA CCTAGGCTCT 240 ATCAGGCTTA GGATTCTTTG AACTTATTTC CACTTTAATT TCTCAGTGGA AGTTAAGAGG 300 30 GGTGAGAAAA CAAAGAAGGG GAAAAACTGA CAACTAACAA AACCAGCACC ACATCGCTAG 360 GTGGTGCTTA CTAATTACCT TCTCAGGATT TTCCTCAGAT TGAAAAGCTT ATGAGGATTT 420 CTTGGGASTC TTAATAACCT GCCTGTTAGT ACAGAGCTTT CCTGATGATA TTTACTCTTG 480 35 AGCACATGIG GTIGITALAAC CITAACITIC TITCTCCAGG ÁGGGIGGIGA TAGAAACAGA 540 TOGTAGTATT TATGAACTGA TETTCTCGTG AAATGTTGAG GGTGGGGAGA AAAGACTTTA 40 AGGGAGGAGA GCCATCTATT TIGTTCCTAA AGCCACCTCT CAGCAGAATC GTCATGTTTT 660 TOTGATECAC COCTOTECTT CATGCCCAAG ATGACTTGCG AGGCAATCTC AGGAGCTGTG 720 GACTTAACCR TIGCAAAGCA CACTGTCTTT CTCAGCGTTC TCTGCAAGTC AGTAGGTGTT 780 45 AGTATGGITG CAAAGTTCAC TGTCTCAGCA AAGTTGAACT GGGCTACCTC TCTACAGCTG TTTCCTCAGA GGGAAAAATC TTGAGACCAG ATGGTGGAGC TCTGGAGTCA GAGGAAATGG 900 50 GTGTCTTCAG CACAAAGCTG CTGCTTTTAC TTCAGCCACT TCTGACATTT TTACATACCG 960 AGCCTGAGAT TRTGTGATTA TCTCAAATCA AATCACTTTG ATGGAGATAA ATAATCAAAA 1020 CTGTTTEATA GTCATTGATT TGGTGAGAAC AGTAATGGAA AATGGTGTTG AAGGACTTCT 1080 55 CATTITIOGA GCTTTCCTTC CAGAGTCCTG GCTGATTGGT GTTCGCTGTT CATCTGAGCC 1140 CCCAAAAGCA TTATTACTGA TACTTGCACA CAGTCAAAAG CGCAGACTGG ATGGATGGTC 1200

	TTTTATAAGG CATTTAAGGG TACACTACTG TGTTTCACTG ACCATACATT TTTCTTAGCC	1260
	CCTCAAGTAA TATAGCACAG AGTTATGAAT GACAATTCCC CTAACCATTC CTCTTCATAT	1320
5	CTGCCTCTTC CCCTTACCAT CGTAATTCTC CAAACTGGTC ATAAAGGCAC TCTGTGAAGA	1380
	TATTGGGGAC TGACATCTTA AGCTCTCACC TGGCTGCAGT AGGAAAGGCC AAACTGACGA	1440
10	CAAAAAAAAA ATTCTTTATA AAGATGATAT GGTAACATGT ATCTTTGCCC TGGGTCTGGG	. 1500
10	TOGGTCCAGT CAGTCTCAGA TTTACAAGCA TTTAGGAGCC TAGGTAAAAG CTGCTAGTAT	1560
	TCTTTTAAAA GTTACATTTA TGACTTGCAA TGATAGAAAA CTCCTTCCAA TTAAATGGCA	1620
15	TTTTATAATA TTATGTGTGT ACTTCACAGT GTTAAAAATA CCCTCATACG TTATTGCATT	1680
	TGATCTTCAC AGAAAGTGCA TTTTAACCAG TACTCTGGGT GCAATAAATA ATATGTAGAA	1740
20	ATTTAAGTCC TCCAATTCCA GCATATCCAG TGAGTTTTGA CAGTGTGTTT ATGTGGAATG	1800
20	TTTAAGGATA TACAATTGTA CTTTATATAA ATTGGTTCTT GTTCTTCTTA AATGTGACAT	1860
	GAAATAATTG TGCTGCTACA TTATACTGGA AATTAACAGG GGAAAAGGGA AGAGCTCTTG	1920
25	GCTCCCTTGA GGTTCTGCTA GTGGTGTTAG GAGTGGTTAC AACTGAGCTT TTAGTAACCA	1980
	TTTAACCGTA TGTAAACTTG GTTTCTAATT AAAAAAAAAT TTCTTTTTCC A	2031

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(2) INFORMATION FOR SEQ ID NO: 33:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 971 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

40

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

	CGCGTCGGAA CTCGGCCGCG GGACATCCAC GGGGCGCGAG TGACACGCGG GAGGGAGAGC	60
	AGTGTTCTGC TGGAGCCGAT GCCAAAAACC ATGCATTTCT TATTCAGATT CATTGTTTTC	120
45	TTTTATCTGT GGGGCCTTTT TACTGCTCAG AGACAAAAGA AAGAGGAGAG CACCGAAGAA	180
	GTGAAAATAG AAGTTTTGCA TCGTCCAGAA AACTGCTCTA AGACAAGCAA GAAGGGAGAC	240
50	CTACTAAATG CCCATTATGA CGGCTACCTG GCTAAAGACG GCTCGAAATT CTACTGCAGC	300
	CGGACACAAA ATGAAGGCCA CCCCAAATGG TTTGTTCTTG GTGTTGGGCA AGTCATAAAA	360
66	OGCCTAGACA TIGCTATGAC AGATATGTGC CCTGGAGAAA AGCGAAAAGT AGTTATACCC	420
55	CCTTCATTTG CATACOGAAA GGAAGGCTAT GCAGAAGGCA AGATTCCACC GGATGCTACA	480
٠	TIGATITITG AGATTGAACT TTATGCTGTG ACCAAAGGAC CACGGAGCAT TGAGACATTT	540
60	AAACAAATAG ACATGGACAA TGACAGGCAG CTCTCTAAAG CCGAGATAAA CCTCTACTTG	600

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	CAAAGGGAAT TIGAAAAAGA TGAGAAGCCA CGTGACAAGT CATATCAGGA TGCAGTTTTA	660
5	GAAGATATTT TTAAGAAGAA TGACCATGAT GGTGATGGCT TCATTTCTCC CAAGGAATAC	720
	AATGTATACC AACACGATGA ACTATAGCAT ATTTGTATTT CTACTTTTTT TTTTTAGCTA	780
	TTTACTGTAC TITATGTATA AAACAAAGTC ACTTTTCTCC AAGTTGTATT TGCTATTTTT	840
0	CCCCTATGAG AAGATATTTT GATCTCCCCA ATACATTGAT TTTGGTATAA TAAATGTGAG	900
	GCTGTTTTGC AAACTTAAAA AAAAAWWAAA AAAACTSGAG GGGGGCCCGT ACCCAANTCG	960
. ~	CCGNATATGA T	971
15		
20	(2) INFORMATION FOR SEQ ID NO: 34: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1792 base pairs	
25	(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:	
30	GAACCCCCTT TCTCCTGGTA AAGGGTAAGG GGGGGGATAA TGTTTACCAC AGGTACGAAA	60
50	TAGTCACTTT AACATTGAGA CCTCTGCCTC ATTGAATTCA GGTTTTTTAA GTACTTGAAA	120
	CTCTTCAGAT TCTCCTTATT TTAGTTTCTT TTTACATTTA TGAAGTAGAA AGCATTGTTT	180
35	TGTAAACTGT TTTGAAAATA AATAGCCTAG TCTCTTATCC TCTTTAGCGT GGATTAAAGG	240
	TGAAGTTCTG CAAATGGGAG AGTGTTCACA GTAGATAGCT CAGATTGATT GAACACATTT	300
40	GAGGAAGAGA CTCCTGCATG AGATACCAGC ATTTTTACAA ATACTTTTTA TGTACATTCT	360
40	TTATITIGIC ATTITIGICAA CCCTCTCCCC AAGCACATCT TCTTTCCTTT TACTATGTCT	420
	ATGTAGGGAA AAACAAAACA AAAAATTGCA CTTACGTTAC ACTCCCAAAA TGTGGGTAAT	480
45	CCGTGTCTTT CAAAAAACAT TTCTGTTTTT TGTTTTGTTT	540
	TGACAAGTTT GGGTGCTTGT GGCACGTATG TATGAAGCGG GAGGGGGATG ASAATTGCCT	600
50	GTCCTTCAGT ARGCTGTAAA AGTAATTTAC ATGTAAGTAA AAAGGGAAAA TAGAATAGAT	660
30	GCCAAAGICA TITATICAGI CCITAGITIT CITATGIGGC ATTACIGCAT CIGCIAGITA	720
	GTGAGAAAGC ACCCTCAGCT TTTACTGCTC CCCTCCCTGC CTGCCAACAC ACTTGATGTG	780
55	TOCANACAGO COTONAGTAT CTGTONGATG ACCTATATAN GGTATTGANT ANGGTATTOT	840
	TGTCAGTTTA GAAATGGACT GGATAAAACT TACTTGGTTG TCATTATTTT ATCTCATTTG	900

TCCTGTTACA TGCCCTATGT TAAGATAATT ATATTGCCAC TAATAATCAA GATGCTAAAT

60

600

		1020
	GAGTATTACA ACTGGCTAAT ATCATTTTT ATATACAAGG GTATGTGTAT ATTTGGAATT	
	GRIATGAGAA ACTCATTIGT ACCCATTIGA GIGATATIGC ACAACAAACA CAGATAYCTA	1080
5	CAGACTCCGT TTTCATTTTC TCGTGTTCTT TATGATAATG ATCTTTGTAG ATTGGTTATT	1140
	TCTGTACTTT ATCTGTAATA AACTTTGTAG ATCCTGTGAA CCATTACTTT GCCTAAATCA	1200
	CTTGAGACTT GAGTCTTTAA TAACAAAGCA TCAATATTCA CTAAAGTCAA TCTCTTTTGA	- 1260
10	GITTCTGTGA CTTGGCTAGA AGCTCTTGAC ACTAAGGGAT TAGTGTTAAT TTTCCCTGGG	1320
	GGTGTTCCAC TAGGGCATTA CTGTATAATG ACTTGATGTT GCCACATAGA CTTCAAGATA	1380
15	TATAATATTT TGAGGATTTT GTTGATTGGC CTATGTTTTA TTGCATAGTG TGAAACGTGT	1440
	AAAGCTTGGT TAACCTGTAT ATAGATAGCT TATTGTTGAC TAGTTATAGT GTATTTAGGG	1500
	TIGCCIGIAA TATITAAGCT TCTTTACTGA TGTGTGTGCT GGTAGGAACA TATAATTTTT	1560
20	GTACATTATA TITACTGAGA TGTTGCCTTT TTTATTTTAC AAATACTTTG GAATTCCAAT	1620
	GIGITITITG CITCCGIGAG GATTAATITG GAAAGGITTT TAATGACATT CCACIGATIT	1680
25	CAGATTITIGC TIGAGATIGA CITCAATAAA TIGICCIGTA TGITCCAAAA AAAAATTAAA	1740
	AAACTCGAGG GGGCCCGGT ACCCAANNCG CCGGATATGA TCGTAAACAA TC	1792
30		
	(2) INFORMATION FOR SEQ ID NO: 35:	
	(i) SEQUENCE CHARACTERISTICS:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid	
35 40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	60
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:	66 120
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT GCCAGCYTCA CYTGCCACYT TYTGCCCCTY TCGGGATGCC TTCGCAGACA GAGYTYTTCG	120
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT GCCAGCYTCA CYTGCCACYT TYTGCCCCTY TCGGGATGCC TTCGCAGACA GAGYTYTTCG CTGCCTGTGG TGGCCAYTCT TTGCTTTTGG TTYTCTTGCC CCTTGGCCTC CCTTTTTGTC	126 186
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT GCCAGCYTCA CYTGCCACYT TYTGCCCCTY TCGGGATGCC TTCGCAGACA GAGYTYTTCG CTGCCTGTGG TGGCCAYTCT TTGCTTTTGG TTYTCTTGCC CCTTGGCCTC CCTTTTTGTC CCCGGGCAGC CTTGTGTGAC CTGCCCTTTT CCCTCCCTTC CTTTCCAGGA CAAGCACGCC	120 180 240
40 45 50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT GCCAGCYTCA CYTGCCACYT TYTGCCCCTY TCGGGATGCC TTCGCAGACA GAGYTYTTCG CTGCCTGTGG TGGCCAYTCT TTGCTTTTGG TTYTCTTGCC CCTTGGCCTC CCTTTTTGTC CCCGGGCAGC CTTGTGTGAC CTGCCCTTTT CCCTCCCTTC CTTTCCAGGA CAAGCACGCC GAGGAGGTGC GGAAAAACAA GGAGCTGAAG GAAGAGGCCT CCAGGTAAAG CCTAGAGGCC	126 186 24 30
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 896 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35: AGTIGNANAC AACAGGACCT GAGTCCTTGG GCAGCACCAG TAGGTTGCCC CYTGCYTCYT GCCAGCYTCA CYTGCCACYT TYTGCCCCTY TCGGGATGCC TTCGCAGACA GAGYTYTTCG CTGCCTGTGG TGGCCAYTCT TTGCTTTTGG TTYTCTTGCC CCTTGGCCTC CCTTTTTGTC CCCGGGCAGC CTTGTGTGAC CTGCCCTTTT CCCTCCCTTC CTTTCCAGGA CAAGCACGCC GAGGAGGTGC GGAAAAACAA GGAGCTGAAG GAAGAGGCCT CCAGGTAAAG CCTAGAGGCC AAAGAACTTT CCAGGTCAGC CGGACAGCTC CAGCAGCTCC ACGTTCCAGG CAGCCTCGMC	120 186 24 30 36

60 AGAACTCAAG GACATTGCAA CCCTGCCCGG CGCAGATCTG ATTTTCACAT CTCTACCTGG

•	ACATTGAGCC	TCCCAGGCAC	CATGTTGAGG	AGAGATGAAA	ACCAGGGCGG	TAGAACTTCA	660
_	GGGTGAAGGA	CAGAGTCCTG	GGTGGGGCAG	CGGCTGCAGG	GCGCACCAGA	GAACCCAGCC	720
5	AGAGGGGGTG	TGAGTACCAG	TGGTGTTGCT	TCCACCCTGC	AGCAGGTGGG	ATGAGGTCTG	780
	TGTGTGTGTG	TGAACCATCA	TTTTTTGATC	ATCATGACCA	ATGAAACATT	GAAAAAAAA	840
10	AAAAAAACTG	GAGGGGGGCC	CGTACCCAAN	TCGCCGNATA	GTGATCGTAA	ACAATC	896

15 (2) INFORMATION FOR SEQ ID NO: 36:

20

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 912 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:

25	TOGACCCACG C	CETCCGGTCA	GCCAGTCGCA	TCCAGCCATG	ACAGCCTTCT	GCTCCCTGCT	60
	CCTGCAAGCG (CAGAGCCTCC	TACCCAGGAC	CATGGCAGCC	CCCCAGGACA	GCCTCAGACC	120
30	AGGGGAGGAA C	GACGAAGGGA	TGCAGCTGCT	ACAGACAAAG	GACTCCATGG	CCAAGGGAGC	180
30	TAGGCCCGGG (CCAKCCGCG	GCAGGGCTCG	CTGGGGTCTG	GCCTACACGC	TGCTGCACAA	240
	CCCAACCCTG (CAGGTCTTCC	GCAAGACGGC	CCTCTTGGGT	GCCAATGGTG	CCCAGCCCTG	300
35	ARGGCAGGGA A	AKGTCAACCC	ACCTGCCCAT	CTGTGCTGAG	GCATGTTCCT	GCCTACCATC	360
	CTCCTCCCTC (CCCCCCTCTC	CTCCCACCAT	CACACCAGCC	ATGCAGCCAG	CAGGTCCTCC	420
40	GGATCACYGT (GGTTKGGTGG	AGGTCTGTCT	GCACTGGGAG	CCTCARGARG	GCTCTGCTCC	480
40	ACCCACTTGG	CTATGGGAGA	GCCAGCAGGG	GTTCTGGAGA	AAAAAACTGG	TGGGTTAGGG	540
	CCTTCGTCCA	GGAGCCAGTT	GAGCCAGGGC	AGCCACATCC	AGGCGTCTCC	CTACCCTGGC	600
45	TCTGCCATCA	GCCTTGAAGG	GCCTCGATGA	AGCCTTCTCT	GGAACCACTC	CAGCCCAGCT	660
	CCACCTCAGC	CTIGGCCTIC	ACGCTGTGGA	AGCAGCCAAG	GCACTTCCTC	ACCCCYTCAG	720
50	CGCCACGGAC	CTYTYTGGGG	AGTGGCCGGA	AAGCTCCCSG	CCTYTCCCC	TGCAGGGCAG	780
50	CCCAAGTCAT	GACTCAGACC	AGGTCCCACA	CTGAGCTGCC	CACACTOGAG	AGCCAGATAT	840
	TTTTGTAGTT	TTTATKCCTT	TGGCTATTAT	GAAAGAGGTT	AGTGTGTTCC	CTGCAATAAA	900
55	CTTGTTCCTG	AG			•		912

^{60 (2)} INFORMATION FOR SEQ ID NO: 37:

297

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1382 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:

	•	
10	AATTCGGCAC GAGCGGAGCC GAGGGAAACT RAGGGCGAAA GITGTGTGTC GTGTTGGCAG	60
	GAGGGCCTAG AAGGGAAAGA CTGTCTAGTG GGACAATGTC ATATTATAAA TTTGGAATGC	120
15	TGAATAGAAA ATTATAGATT TTGATATTGA AGGAAATGAA GCGAAGCYTA AATGAAAATT	180
	CAGCTCGAAG TACAGCAGGC TGTTTGCCTG TTCCGTTGTT CAATCAGAAA AAGAGGAACA	240
	GACAGCCATT AACTTCTAAT CCACTTAAAG ATGATTCAGG TATCAGTACC CCTTCTGACA	300
20	ATTATGATTT TCCTCCTCTA CCTACAGATT GGGCCTGGGA AGCTGTGAAT CCAGAGTTKG	360
	CTCCTGTAAT GAAAACAGTG GACACCGGGC AAATACCACA TTCAGTTTCT CGTCCTCTGA	420
25	GAAGTCAAGA TTCTGTCTTT AACTCTATTC AATCAAATAC TGGAAGAAGC CAGGGTGGTT	480
	GGAGCTACAG AGATGGTAAC AAAAATACCA GCTTGAAAAC TTGGRATAAA AATGATTTTA	540
	AGCCTCAATG TAAACGAACA AACTTAGTGG CAAATGATGG AAAAAATTCT TGTCCAATGA	600
30	GTTCGGGAGC TCAACAACAA AAACAATTAA GAACACCTGA ACCTCCTAAC TTATCTCGCA	660
	ACAAAGAAAC CGAGCTACTC AGACAAACAC ATTCATCAAA AATATCTGGC TGCACAATGA	720
	GAGGGCTAGA CAAAAACAGT GCACTACAGA CACTTAAGCC CAATTTTCAA CAAAATCAAT	780
35	ATAAGANACA AATGTTGGAT GATATTCCAG AAGACAACAC CCTGAAGGAA ACCTCATTGT	840
	ATCAGTTACA GITTAAGGAA AAAGCTAGTT CTTTAAGAAT TATTTCTGCA GITATTGAAA	900
40	GCATGAAGTA TIGGCGIGAA CATGCACAGA AAACTGTACT TCTTTTTGAA GTATTAGCIG	960
	TTCTTGATTC ACCTGTTACA CCTGGCCCAT ATTATTCGAA GACTTTTCTT ATGAGGGATG	1020
45	GGAAAAATAC TCTGCCTTGT GTCTTTTATG AAATCGATCG TGAACTTCCG AGACTGATTA	1080
45	GAGGCCGAGT TCATAGATGT GTTGGCAACT ATGACCAGAA AAAGAACATT TTCCAATGTG	1140
	TTTCTGTCAG ACCGCCGTCT GTTTCTGAGC AAAAAACTTT CCAGGCATTT GTCAAAATTG	1200
50	CAGATGTTGA GATGCAGTAT TATATTAATG TGATGAATGA AACTTAAGTA GTGATAAAAG	1260
	GAAGTTTAGC ATAAATTATA GCAGTTTTCT GTTATTGCTT AATTTACCAT CTCCATAGTT	1320
	TTATAGCTAC TATTGTATTT CACTTGTTGA ATTAAAGTAT TTGAATTCTT TTAAAAAAAA	1380
55	aa	1382

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(2) INFORMATION FOR SEQ ID NO: 38:

5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 872 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:	
10	GGGCTACTIC AAAGCCCTGG GCCTTATTIC TICAGGTAAA AAAATATAAA GTCAGATCTC	60
	ATCCCGGCTG GCCATGCTGT TAGACCCTTT CATCCTTCTC TTCTGCCTCT TCTCAACAGC	120
15	TGCCCAGTCC TGTTTGGAAT TCATATACAT ACAGTTCTAA TACTGATGTA TTTACCCTCA	180
	TAAGCCACTC AACCCAGAAT CTTATTTGAA TTATAATCCA GAAACATCAG GTGACGTGTG	240
	AGACTACTGT ATGAGAAAGA GACAGTTTAA GGGTCAGTCC AATGGAAAAA AGAGTTCTCA	300
20	GAGCTITCTT TAGCTTATTC TCATCAAAGA GCTTTCTCTG CAGAAGGAAC CTACTGGTTC	360
	CTCCTTTCCA GTCCTAGAAA TCCTGACCTA GAGTGGCTTA ATCCTGCTAG CACCTCTCTC	420
25	TOGCACTOTG GTGCCAAATG ACTOCAGGAA CTGGGCCATG ATGTGGTGGG AATGACCTTA	480
	CCCTGAGCAT GTCACTCATG CATTGAACAA CAGCTAAGAG CAGAGCTTAG AGCTTAGAGC	540
	TGGGCCCTGT AAGGTGAGAG GAATCACATC CTGCAGAAGT CTGTCCTGAG AAGCAGGTAC	600
30	TCCTGTCACA GCAGAGACAC AGTGGATACC TGAGTAACAA TAATACAAGA CAGGACGTGG	660
	GMACAGCAAA AGATTTGGGT GTCAGAAGAR GCCGAGAACA CTTYCAGGCA GGAACATTCA	720
35	RARTTGTTCT TGGAGGAART AGGCMCSAAG GCTGGGCAGG ATTTCMCGGG GCAGAGATGG	780
	AGCAAGCAAT TGAAATGAAA GCCATGGCAT GGGAAAAGGA GCACTGGCCA CAGGGAGTGC	840
40	AACGTTGTGA TGCAAGGCCA CTGTGGAGCC AT	872
	(2) INFORMATION FOR SEQ ID NO: 39:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 812 base pairs (B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:	
		60
55	GGCAGAGGCT CACCCCAGCA GAGATTGAGG GGGAACCGTG ATGAAATTTT TAAGTATTCT	120
	GCTTGATGAT AATAATTITY CTCTTATGIT AATGITGGCT CCGTTTGGGT GITTAGCTTT	180
	TGAAAGGAGT ATGAAAATGC GGAATGGGGC TTTGGGGCTT GAGGAGGTGT GATCTCTAGT	
60	CTITAAAAAA TITAATIGCA CAAATAGAAA TAATICACCC ACATTATIGA ACCCCACTAA	240

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	AGCATATOCT TITTGTCCAT ATTCCTTTCC TGCTGCCCTC GTGTGTACCA TTATTACTCA	300
_	GTTGTGATTT GAGCTCGTTC CACTTAAAGT CATTCATAGA TACTTTTGCG TCGTGTTKGA	360
5	ATATTTATTG AATTTCTATT CTGTGTTTTA CTTAATTACT TTATTATGGA ACCTTTACAC	420
	AGGTCTGGTG TACTTGTTCT TTGAAAAGTC TTATGTTGAC CACCATCACT GAGCATATAG	480
10	CTTTTTCCTT ATTTCCTTGG GATAATTACC CGAAGTGGAA ATACCGAATC AAACTTCTGT	540
	TTTCTTTCTT TGGCACTATT ATATAAATTG TTTTCCAAAC AAGGCATGTT TACAATAGAC	600
1 5	ATTTTTCAAA ATCTGGGTAT TTGTCCTATT TTGCTCTCTG TATGCAGAAT TCAGCGGGGT	660
15	GCCAAGTCGT TTTCTGTGTG GGTTGAGAGA CAGGCTGTGC AGCCCACTGT TGCATAGGAC	720
	TAACTACTAC AAATCATGCT GAGACCGAGC TATTTTTGCT GCTTAGARGC TTTGCAGCCT	780
20	TGAGTAAGTT TCGNCATCTG GAAACNITGN AA	812
•		
25	(2) INFORMATION FOR SEQ ID NO: 40:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1515 base pairs	
20	(B) TYPE: nucleic acid	
30	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:	
35	AATTCGGCAC GAGGGAAATT CAAGCACTTT TCCTAAAAGA AGGGGGAATG GATGCTGAAA	60
	CAACACGINI CCCACAAAGG GAGCAGACAC TGGGCTTGTG AAGCTGCCCC ATACCTTCCC	120
40	CACAGAACTG GGGTCCGGCC TCCCTGACAT GCAGATTTCC ACCCAGAAGA CAGAGAAGGA	180
40	GCCAGTGGTC ATGGAATGGG CTGGGGTCAA AGACTGGGTG CCTGGGAGCT GAGGCAGCCA	240
	COGTTTCAGC CTGGCCAGCC CTCTGGACCC CGAGGTTGGA CCCTACTGTG ACACACCTAC	300
45	CATGOGGACA CTCTTCAACC TCCTCTGGCT TGCCCTGGCC TGCAGCCCTG TTCACACTAC	360
	CCTGTCAAAG TCAGATGCCA AAAAAGCCGC CTCAAAGACG CTGCTGGAGA AGAGTCAGTT	420
50	TTCAGATAAG CCGGTGCAAG ACCGGGTTT GGTGGTGACG GACCTCAAAG CTGAGAGTGT	480
50	GETTCTTGAG CATCGCAGCT ACTGCTCGGC AAAGGCCCGG GACAGACACT TTGCTGGGGA	540
	TGTACTGGGC TATGTCACTC CATGGAACAG CCATGGCTAC GATGTCACCA AGGTCTTTGG	600
55	CACCAACTIC ACACACTCT CACCCCTCTG CCTGCACCTG AAGAGACGTG GCCGTGAGAT	660

CTTTGAGGTC ACGGCCTCC ACGACGTGGA CCAAGGGTGG ATGCGAGCTG TCAGGAAGCA

TGCCAAGGGC CTGCACATAG TGCCTCGGCT CCTGTTTGAG GACTGGACTT ACGATGATTT

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	CCGGAACGTC TTAGACAGTG AGGATGAGAT AGAGGAGCTG AGCAAGACCG TGGTCCAGGT	840
	GGCAAAGAAC CAGCATTTCG ATGGCTTCGT GGTGGAGGTC TGGAACCAGC TGCTAAGCCA	900
5	GAAGCGCCTG ACCGACCAGC TGGGCATGTT CACGCACAAG GAGTTTGAGC AGCTGGCCCC	960
	CGTGCTGGAT GGTTTCAGCC TCATGACCTA CGACTACTCT ACAGCGCATC AGCCTGGCCC	1020
10	TAATGCACCC CTGTCCTGGG TTCGAGCCTG CGTCCAGGTC CTGGACCCGA AGTCCAAGTG	1086
10	GCGAAGCAAA ATCCTCCTGG GGCTCAACTT CTATGGTATG GACTACGCGA CCTCCAAGGA	1140
	TGCCCGTGAG CCTGTTGTCG GGGCCAGGTA CATCCAGACA CTGAAGGACC ACAGGCCCCG	120
15	GATGGTGTGG GACAGCCAGG YCTCAGAGCA CTTCTTCGAG TACAAGAAGA GCCGCAGTGG	126
	GAGGCACGTC GTCTTCTACC CAACCCTGAA GTCCCTGCAG GTGCGGCTGG AGCTGGCCCG	132
20	GGAGCTGGGC GTTGGGGTCT CTATCTGGGA GCTGGGCCAG GGCCTGGACT ACTTCTACGA	138
20	CCTGCTCTAG GTGGGCATTG CGGCCTCCGC GGTGGACGTG TTCTTTTCTA AGCCATGGAG	144
	TGAGTGAGCA GGTGTGAAAT ACAGGCCTTC ACTCCGTTAA AAAAAAAAAA	150
25	AAAAAAAAA AAAAA	151

30 (2) INFORMATION FOR SEQ ID NO: 41:

35

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 704 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:

40	AAGATGGTGG	CGCCCAGAGC	TTCGCTCTAT	OCTGCTCCCC	TGAGAGAGGC	GTTTCCATCA	60
	ACCAGTTTTG	CAAGGAGTTC	AATGAGAGGA	CAAAGGACAT	CAAGGAAGGC	ATTCCTCTGC	120
45	CTACCAAGAT	TTTAGTGAAG	CCTGACAGGA	CATTTGAAAT	TAAGATTGGA	CAGCCCACTG	180
73	TTTCCTACTT	CCTGAAGGCA	GCAGCTGGGA	TTGAAAAGGG	GGCCCGGCAA	ACAGGGAAAG	240
	AGGTGGCAGG	CCTGGTGACC	TTGAAGCATG	TGTATGAGAT	TGCCCGCATC	AAAGCTCAGG	300
50	ATGAGGCATT	TGCCCTGCAG	GATGTACCCC	TGTCGTCTGT	TGTCCGCTCC	ATCATCGGGT	360
	CICCCCCITC	TCTGGGCATT	CCCCTCCTGA	AGGACCTCAG	TTCAGAAGAG	CTTGCAGCTT	420
55	TCCAGAAGGA	ACGAGCCATC	TTCCTGGCTG	CTCAGAAGGA	GGCAGATTTG	GCTGCCCAAG	480
33	AAGAAGCTGC	CAAGAAGTGA	CCCTTGCCCC	ACCAACTCCC	AGATTTCAAA	GGAGGTAGTT	540
	GCAAAAGCTG	TGCCCAAGGG	GAGGAAGGAG	GTCACACCAA	TATGATGATG	GTTTTCATGA	600
60	CTTTGAATGA	TATATTTTTG	TACATCTAGC	TGTATCGAGG	CATCAGGCCT	GAATAAACAT	660

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•	CCTTTCTTAA AAAAAAAAAA AAAAAAAAAA AAAAAAAA	704
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	(2) INFORMATION FOR SEQ ID NO: 42:	
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1094 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	•
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:	
	GGCAGCTITC TTACAAACCC ATCCTTCTGA AATGTTGCTT CAAATTCATC CTCTGCTCCC	60
••	CAGTCCCACT ATTCCACACA TACTGTTACT GITTCTTTAT CCTACTTTCT CAATTTTGGA	120
20	ACATAGITIGC AGITACTICA TIGAATACCT GIGGGITTIGC CIGTIGITCT GICTGTCTCT	180
	GTGGTTCTTG TAATANTGGA TCCCAGAGAT AAAATGGACA GTTGTNATGC ACAGTTAATT	240
25	CAGAAACTAG ACCTTACTTG CTGTGTGAAA TACCAACTAA ATTCTCAGTG AACTCAGCTG	300
	ANCTITATCT CCTTTTGTTT CCCCAATTTA TAATTTCAGT TCAGGCCCAG AAAGATGGAA	360
	TCCCAGCTAA GAAATACAAG TTACACCCTG TACTAGCAGC CCATGTGTGC ATGTTCTTTA	420
30	AGTGCTCTTG CAGCTATGTC ATTTATATTG ATTTCCCTGT ATTATTATAA GCAAAGCAAA	480
	TTTGAGGAAA AAAACCCATA ATACCACACC TCATTTTTTT CAAGTAATAG GGTCATAAGT	540
35	CTCATYCTYC ATATAATATG TTGAGTATGC AGTATATTAT GTGTTAGGCT CTGGANAGGC	600
	AGAGGITAGA TCATGIWACA GATÇATATCK GATTAGGCAG ATAAACAGTA TTTTAACCIT	660
	TICCTTATTA TATGTAACIT GCTTTCAGGT TTTTTAATGT TACTATTATG TCTTTAATAT	720
40	ATTATOTTA TITGIACTIT TGIATACAGA GIGATTITCC TITTITAAAA AAAATTGIGT	780
٠.	CTTTAGGATG GATTCCAAAG ATGTGGAATC AGTAGGTTTA AGGAATATGG ATATTTTGGC	840
45	TGGCAAGGTG GCTCACACCT GTAATCCCAG CACTTTGGGA GGCTGAGGTG GGTGGATCAC	900
	CTGAAGTCAG GAGTTCGAGA CCAGCCTGAC CAACATGGCG AAACCCTGTT TNTACTAAAG	960
	ACACACWWAA AATTRGCCAG TGGTGGTGGC ATGTGCTTGT AGTCCCACTT AGCTACTCGA	1020
50	GAGGCTGAGG CAGGAGAATC GCTTGAACCC GGGAGGCAGA GGTTGCAGTG AGGCAAGATG	1080
	GCACCICTAC ACTC	1094

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(2) INFORMATION FOR SEQ ID NO: 43:

60 (i) SEQUENCE CHARACTERISTICS:

302

(A) LEWIH: 1321 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:

	TOGOTTREGO CATCACCOTT COOTTGGCTG GAACTACTGG ACAGACCCTT TTGAGATGTG	60
10	CONGREGATE PERCENCIAGE TOTAL TOTAL TOTAL TOTAL	120
	TIGISTAGIC TRACCIGIAT GCIGAAATIG GGCGIGIGIT GGAGGGCTIC TIAGCICTIT	180
	GOTGAGATIG TATTICTATG TGTTTGTATC ASCIGAATGT TGCTGGAAAT AAAACCTTGG	240
15	TITGINAAGG CICYPPPING TGGGAAGIAA GIAGGGGAAA AGGICTITGA GGGIICCTAG	300
	GCTCCTTTGT ACAACAGGAA AATGCCTCAA AGCCTTGCTT CCCAGCAACC TGGGGCTGGT	360
20	TOCCAGTGCC TGGTCCTGCC CCTTCCTGGT TCTTATCTCA AGGCAGAGCT TCTGAATTTC	420
	AGGCCTTCAT TCCAGAGCCC TCTTGTGGCC AGGCCTTCCT TTGCTGGAGG AAGGTACACA	480
25	GGGTGAAGCT GATGCTGTAC TTGGGGGATC TCCTTGGCCT GTTCCACCAA GTGAGAGAAG	540
23	GTACTTACTC TTGTACCTCC TGTTCAGCCA GGTGCATTAA CAGACCTCCC TACAGCTGTA	600
	GGAACTACTG TCCCAGASCT GAGGCAAGGG GATTTCTCAG GTCATTTGGA GAACAAGTGC	660
30	TITAGTAGTA GETTAAAGTA GEAACTGCTA CEGETATETAG EGGGETGGAA ETCAGAAGAA	720
	ATTTEAAGAC CAGATCATEG STEGTCTECA TETGAATGAA CAGGAATGAG CCGGACAGCC	780
35	TEGCTETCAT TECTTTCTTC CTCCCCATTT GGACCCTTCT CTGCCCTTAC ATTTTTGTTT	840
	CTCCATCTAC CACCATCCAC CAGTCTATTT ATTAACTTAG CAAGAGGACA AGTAAAGGGC	900
	CCTCTTGGCT TGATTTTGCT TCTTTCTTTC TGTGGAGGAT ATACTAAGTG CGACTTTGCC	960
40	CTATCCTATT TOGRAATCCC TAACAGAATT GAGTTTTCTA TTAAGGATCC AAAAAGAAAA	1020
٠.	ACALANTECT ANTERAGCCA TCAGTCAAGG GTCACATGCC AATAAACAAT AAATTTTCCA	1080
45	GAAGAAATGA AATSCAACTA GACAAATAAA GTAGAGCTTA TGAAATGGTT CAGTAAGGAT	1140
45	GAGLITGIIG TITLITGIIT TGITTIGIIT TGKTITITTA AAGACGGAGT CTCGCTCTGT	1200
	CACTCAGGCT GGAGTGCAGT GGTATGATCT TGGCTCACTG TAACCTCCGC CTCCCGGGTT	1260
50	CAAGCCATTC TOCTGCCTCA GTCTCCTGAG TAGCTGGGAT TACAGGTGCG TGCCACCATG	1320
	CCTGGCT:AT TITIGTGTTT TTAGTAGAGA CAGGGTTTCA CCATGTTGGT CGGGCTGGTC	1380
55	TCAAACTECT GACCTCTTGA TCCGCCTGCC TTGGCCTCCC AAAGTGATGG GATTACAGAT	1440
<i></i>	GTGACCCACC CGTSCCCTAG CCAAGGATGA GATTTTTAAA GTATGTTTCA GTTCTGTGTC	1500
	ATGGTTGGAA GACAGAGTAG GAAGGATATG GAAAAGGTCA TGGGGAAGCA GAGGTGATTC	1560
60	ATGGCTCTGT GAATTTGAGG TGAATGGTTC CTTATTGTCT AGGCCACTTG TGAAGAATAT	1620

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5	GAGTCAGTTA	TTGCCAGCCT	TOGAATITAC	TTCTCTAGCT	TACAATGGAC	CTTTTGAACT	1686
	GGAAAACACC	TIGICIGCAT	TCACTTTAAA	ATGTCAAAAC	TAATTTTTAT	AATAAATGTT	1740
	TATTTCACA	TTGAAAAAA	AAAAAAATTT	AAAAACYCGG	GGGGGGCCCS	GHACCCCATT	180
	NGCCCCTAAG	GGGGGGGTT	T				182

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(2) INFORMATION FOR SEQ ID NO: 44:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1024 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:

GCCCCACAGT TGAAGAACCG ACCGAGGGAC TGGGAGTCGT TAGTGAGGAT GACGCGGCAT 60 GOCAAGAACT GCACCGCAGG GCCGTCTACA CCTACCACGA GAAGAAGAAG GACACAGCGG 120 CCTCGGGCTA TGGGACCCAG AACATTCGAC TGAGCCGGGA TGCCGTGAAG GACTTCGACT 180 GCTGTTGTCT CTCCCTGCAG CCTTGCCACG ATCCTGTTGT CACCCCAGAT GGCTACCTGT 240 ATGAGCGTGA GGCCATCCTG GAGTACATTC TGCACCAGAA GAAGGAGATT GCCCGGCAGA 300 TGAAGGCCTA CGAGAAGCAG CGGGGCACCC GGCGCGAGGA GCAGAAGGAG CTTCAGCGGG 360 CGGCCTCGCA GGACCATGTG CGGGGCTTCC TGGAGAAGGA GTCGGCTATC GTGAGCCGGC CCCTCAACCC TTTCACAGCC AAGGCCCTCT CGGGCACCAG CCCAGATGAT GTCCAACCTG 480 COCCCAGTGT GGGTCCTCCA AGTAAGGACA AGGACAAAGT GCTGCCCAGC TTCTGGATCC 600 CGTCGCTGAC GCCCGAAGCC AAGGCCACCA AGCTGGAGAA GCCGTCCCGC ACGGTGACCT GCCCCATGTC AGGGAAGCCC CTGCGCATGT CGGACCTGAC GCCCGTGCAC TTCACACCGC TAGACAGCTC CGTGGACCGC GTGGGGCTCA TCACCCGCAG CGAGCGCTAC GTGTGTGCCG 720 TGACCCGCGA CAGCCTGAGC AACGCCACCC CCTGCGCTGT GCTGCGGCCC TCTGGGGCTG 780 TOGTCACCCT CGAATGCGTG GAGAAGCTGA TTCGGAAGGA CATGGTGGAC CCTGTGACTG 840 GAGACAAACT CACAGACCGC GACATCATCG TGCTGCAGCG GGGCGGTACC GSTTCGCGGG 900 960 CTCCGGAGTG AAGCTGCAAG CGGAGAAATC ACGGCCGGTG ATGCAGGCCT GAGTGTGTGC 1020 1024 AAAA

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(2) INFORMATION FOR SEQ ID NO: 4	(2)	INFORMATION	FOR	SEO	ID	NO:	45
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5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 983 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:	
	CGACACGCT GCGAGAAGAC GACAGAAGGG CCCGACCGCG AGCCGTCCAG GTCTCAGTGC	60
15	TGTGCCCCCC CCAGAGCCTA GAGGATGTTT CATGGGATCC CAGCCACGCC GGGCATAGGA	120
13	GCCCCTGGGA ACAAGCCGGA GCTGTATGAG GAAGTGAAGT	180
	AGGGAGAAGT ACGACAACAT GGCAGAGCTG TTTGCGGTGG TGAAGACAAT GCAAGCCCTG	240
20	GAGAAGGCCT ACATCAAGGA CTGTGTCTCC CCCAGCGAGT ACACTGCAGC CTGCTCCCGG	300
	CTCCTGGTCC AATACAAAGC TGCCTTCAGG CAGGTCCAGG GCTCAGAAAT CAGCTCTATT	360
25	GACGAATTCT GCCGCAAGTT CCGCCTGGAC TGCCCGCTGG CCATGGAGCG GATCAAGGAG	420
25	GACCGCCCCA TCACCATCAA GGACGACAAG GGCAACCTCA ACCGCTGCAT CGCAGACGTG	480
	GTCTCGCTCT TCATCACGGT CATGGACAAG CTGCGCCTGG AGATCCGCGC CATGGATGAG	540
30	ATCCAGCCCG ACCTGCGAGA GCTGATGGAG ACCATGCACC GCATGAGCCA CCTCCCACCC	600
	GACTTTGAGG GCCGCCAGAC GGTCAGCCAG TGGCTGCAGA CCCTGAGCGG CATGTCGGCG	660
25	TCAGATGAGC TGGACGACTC ACAGGTGCGT CAGATGCTGT TCGACCTGGA GTCAGCCTAC	720
35	AACGCCTTCA ACCGCTTCCT GCATGCCTGA GCCCGGGGCA CTAGCCCTTG CACAGAAGGG	780
	CAGAGTOTGA GGCGATGGCT CCTGGTCCCC TGTCCGCCAC ACAGGCCGTG GTCATCCACA	840
40	CAACTCACTG TCTGCAGCTG CCTGTCTGGT GTCTGTCTTT GGTGTCAGAA CTTTTGGGCC	900
	GGGCCCCTCC CCACAATAAA GATGCTCTCC GACCTTCAAA AAAAAAAAAA	960
45	KGSGGCCGGT CCCCANTCCC CCC	983
45		
50	(2) INFORMATION FOR SEQ ID NO: 46:	
50	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 2421 base pairs (B) TYPE: nucleic acid	
55	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:	

CCGGCTGATC GCTGCCGCTC CGCCAATACA ATAGAGCCAK CCACTACCAG CAGCCTGGCC

•	CICITCCTCC	TTCTCCAGAG	AGACCAATCC	AGCCGAACTC	GGGGTTTGCC	TGAGGAGAAG	120
	GAGGAAGTGA	CCATGGACAC	AAGTGAAAAC	AGACCTGAAA	ATGATGTTCC	AGAACCTCCC	180
5	ATGCCTATTG	CAGACCAAGT	CAGCAATGAT	GACCGCCCGG	AGGGCAGTGT	TGAAGATGAG	240
	GAGAAGAAAG	AGAGCTCGCT	GCCCAAATCA	TTCAAGAGGA	AGATCTCCGT	TGTCTCAGCT	300
10	ACCAAGGGGG	TGCCAGCTGG	AAACAGTGAC	ACAGAGGGGG	GCCAGCCTGG	TCGGAAACGA	. 360
10	CGCTGGGGAG	CCAGCACAGC	CACCACACAG	AAGAAACCTT	CCATCAGTAT	CACCACTGAA	420
	TCACTAAAGA	GCCTCATCCC	CGACATCAAA	CCCCTGGCGG	GGCAGGAGGC	TGTTGTGGAT	480
15	CTTCATGCTG	ATGACTCTCG	CATCTCTGAG	GATGAGACAG	AGCGTAATGG	CGATGATGGG	540
	ACCCATGACA	AGGGGCTGAA	AATATGCCGG	ACAGTCACTC	AGGTAGTACC	TGCAGAGGGC	600
20	CAGGAGAATG	GGCAGAGGGA	AGAAGAGGAA	GAAGAGAAGG	AACCTGAAGC	AGAACCTCCT	660
20	GTACCTCCCC	AGGTGTCAGT	AGAGGTGGCC	TTGCCCCCAC	CTGCAGAGCA	TGAAGTAAAG	720
	AAAGTGACTT	TAGGAGATAC	CTTAACTCGA	CGTTCCATTA	GCCAGCAGAA	GTCCGGAGTT	780
25	TCCATTACCA	TTGATGACCC	AGTCCGAACT	GCCCAGGTGC	CCTCCCCACC	CCGGGGCAAG	840
	ATTAGCAACA	TTGTCCATAT	CTCCAATTTG	GTCCGTCCTT	TCACTTTAGG	CCAGCTAAAG	900
30	GAGTTGTTGG	GCCCACAGG	AACCTTGGTG	GAAGAGGCCT	TCTGGATTGA	CAAGATCAAA	960
30	TCTCATTGCT	TTGTAACGTA	CTCAACAGTA	GAGGAAGCTG	TTGCCACCCG	CACAGCTCTG	1020
	CACGGGGTCA	AATGGCCCCA	GTCCAATCCC	AAATTCCTTT	GTGCTGACTA	TGCCGAGCAA	1080
35	GATGAGCTGG	ATTATCACCG	AGGCCTCTTG	GTGGACCGTC	CCTCTGAAAC	TAAGACAGAG	1140
	GAGCAGGGAA	TACCACGGCC	CCTGCACCCC	CCACCCCCAC	CCCCGGTCCA	GCCACCACAG	. 1200
40	CACCCCCGGG	CAGAGCAGCG	GGAGCAGGAA	CGGGCAGTGC	GGGAACAGTG	GGCAGAACGG	1260
40	GAACGGGAAA	TGGAGCGGCG	GGAGCGGACT	CGATCAGAGC	GTGAATGGGA	TCGGGACAAA	1320
٠.	GTTCGAGAAG	GGCCCCGTTC	CCGATCAAGG	TCCCGTRACC	GCCGCCGCAA	GGAACGTGCG	1380
45	AAGTCTAAAG	AAAAGAAGAG	TGAGAAGAAA	GAGAAAGCCC	AGGAGGAACC	ACCTGCCAAG	1440
	CTGCTGGATG	ACCTTTTCCG	AAAGACCAAG	GCAGCTCCCT	GCATCTATTG	GCTCCCACTG	1500
50	ACTGACAGCC	AGATCGTTCA	GAAAGAGGCA	GAGCGGGCCG	AACGGCCAA	GGAGCGGGAG	1560
50	AAGCGGCGAA	AGGAGCAAGA	AGAAGAAGAG	CAAAAGGAGC	GGGAGAAGGA	AGCCGAGCGG	1620
	GAACGGAACC	GACAGCTGGA	GCGAGAGAAA	CGTCGGGAGC	ACAGTCGGGA	GAGGGACAGG	1680
55	GAGAGAGAGA	GAGAAAGGGA	GCGGGACAGG	GGGGACCGAG	ATCCCGATAC	GGAAAGGGAC	1740
	CGAGAACGAG	GCAGGGAAAG	GGATCGCAGG	GACACCAAGO	GCCACAGCAG	AAGCCGGAGT	1800
60	CGGAGCACAC	CTGTGCGGG	CCGGGGTGGG	CGCCGCTAGC	TGGGAAAACA	CTAGACCTCC	1860

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•	AGGTACCAGC CACTO	GGCCC CAGGGGGTT	A TGGCCACAGA	GGGATAGGCA	CAGTCTCCAC	1920
	CACCCTGGAG CCAAG	GGTCT TTCACATCA	C CTATCCCTAC	ATACATACCA	AATGGAAAAG	1930
5	TGGCCATCCT TTTCC	CCCCA AACACACCC	C CTTAACCTAT	CTCTTGGGAC	TTAGCCCGAC	2340
	CCTCCCTCTC ATTTC	CCATT AAGTCTGAG	A GGCAAGAGCT	AGGTTAGGCA	AGGAGGTGGT	2100
10	TOGCCAGAGA TOGGC	GAACAG CCAGGTGCC	C CAGTCCTCTG	ATTITICCIC	CATCCTGCTT	2160
10	ACCACCTCCC TGGGT	PACTTA CAGCCTTCT	C TTGGGAACAG	CCGGGGCCAG	GACTGGGTCA	2220
	CCTATGAGCT GAATC	CAGCAT CTCCTCCTC	A GTCCCAGGGC	CCCTGCAGTT	CCCAGTCTCT	2280
15	TCTGTCCTGC AGCCC	CTTGCC TCTTTCCC	C AGGITCCACT	TATATATCCAC	CTTTTCCTTT	2340
	TGTTCAATTT TTAT	TTTTAT TTTTTTAT	T ATTAAATGAT	GTGGTCTATG	GAAAAAAAA	2400
20	TAAAAATCTG ACTT	AGTTTT A				2421

(2) INFORMATION FOR SEQ ID NO: 47:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 840 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:

50 CTCAAACTCC TGAGCTGAAG CGATCTACCT GCCTCAGCTA GGATTACAGG TGTGAGCCAC CGCACCCAAC CTCAATAAGC KTATTTGATA AAAKATATGC AAGCTCCCTT TATKCACTTT 120 TCATTCAGAA TGTTTAGTAA TTTGTATTGT TTTTCAGATT TTCAGCCCAA TATATCTCCY 180 TOCCCACTOT GTCACTGTAT TCTACCTAWA CATCATCACG TGTTTCTGCT ATTGGCTGTA 240 TGATGGAACA CTGCGGCTCA TTTTCCTGAA AACTGCCGAT AGTGCATAGA RTGCTGGGAT 300 GGAAACCAGA ARCTITGAAT TCAAGCCTTG GTTCTGCCTT GTTTTTGCTT GGGTGGCCTT 360 420 GAGTCAGCCA CATACCTTTT AAAATCTCAA TTTATTAGAA ATTATTCCAA ATCAAAATCA AATGAGAAGG TATATACAAA AGTGCTTTAT CCCACAATAA ACTATTCAAG AGAGAGCAAA 480 GGAGAGGACA TITACTCAAC ACCTCCTAAA AGGCAGCCAG TGAAATTAGG CATTITATTT 540 AATCCTCCTG GCAACTCTGA GAGTAAAGCA TTATTAATCC CATTTTGGCT GTTTAAAGAA 500 560 ATTATTTGCA CTAGATTCCA GCTGTAGTTT AGYTTCAGAA AAAAAAATCC TGAGATGTGA ATTCACAGCT TTCTGGGTTT AAAGCCCAAG CTCTATCACA TCATGCTATT ATTGTTACAT 720 TACTOCTAGT TCTATGAAAA GAAATACTAA TTTATGAAAT ACATCTTATC CAAAAAAAAA 780 AAAAAAAAC TGGGAGGGG GGCCCGTACC CAAATCGCCG GATAGTGATC GTAAACAATC 340 PCT/US98/11422

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5	(2)	INFORMATION	FOR	SEO	ID	NO:	48:
.)	(2)	TUL OKUMI TON	FUR	SEQ		110.	

WO 98/54963

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2432 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:

15	GGCACGAGGC CCGGAACGCT GAGGAAGGCC CCGTCCCGCC TTCCCCGGCG CGCCATGGAG	60
	CCCCGGGCGG TTGCAGAAGC CGTGAGAACG GGTGAGGAGG ATGTGATTAT GGAAGCTCTG	120
20	CGGTCATACA ACCAGGAGCA CTCCCAGAGC TTCACGTTTG ATGATGCCCA ACAGGAGGAC	180
20	CGGAAGAGAC TGGCGGASTG CTGGTCTCCG TCCTGGAACA GGGCTTGCCA CCCTCCCACC	240
	GTGTCATCTG GCTGCAGAGT GTCCGAATCC TGTCCCGGGA CCGCAACTGC CTGGACCCGT	300
25	TCACCAGCCG CCAGAGCCTG CAGGCAYTAG CCTGYTATGY TGACATCTCT GTCTCTGAGG	360
	GGTCCGTCCC AGAGTCCGCA GACATGGATG TTGTACTGGA GTCCCTCAAG TGCCTGTGCA	420
20	ACCTOGTGCT CAGCAGCCCT GTGGCACAGA TGCTGGCAGC AGAGGCCCGC CTAGTGGTGA	480
30	AGCTCACAGA GCGTGTGGGG CTGTACCGTG AGAGGAGCTT CCCCCACGAT GTCCAGTTCT	540
	TTGACTTGCG GCTCCTCTTC CTGCTAACGG CACTCCGCAC CGATGTGCGC CANAGCTGTT	600
35	TCAGGAGCTG AAAGGAGTGC GCCTGCTAAC TGACACACTG GAGCTGACGC TGGGGGTGAC	660
	TCCTGAAGGG AACCCCCCAC CCACGCTCCT TCCTTCCCAA GAGACTGAGC GGGCCATGGA	720
	GATCCTCAAA GTGCTCTTCA ACATCACCCT GGACTCCATC AAGGGGGAGG TGGACGAGGA	780
40	AGACGCTGCC CTTTACCGAC ACCTGGGGAC CCTTCTCCGG CACTGTGTGA TGATCGCTAC	840
٠.	TOCTOGAGAC COCACAGAGG AGTTCCACGG CCACGCAGTA ASCCTCCTGG GGAACTTGCC	900
45	CCTCAAGTGT CTGGATGTTC TCCTCACCCT GGAGCCACAT GGAGACTCCA CGGAGTTCAT	960
	GGGAGTGAAT ATGGATGTGA TTCGTGCCCT CCTCATCTTC CTAGAGAAGC GTTTGCACAA	1020
	GACACACAGG CTGAAGGAGA GTGTAGCTCC CGTGCTGAGC GTGCTGACTG AATGTGCCCG	1080
50	GATGCACCGC CCAGCCAGGA AGTTCCTGAA GGCCCAGGTG CTGCCCCCTC TGCGGGATGT	1140
	GAGGACACGG CCTGAGGTTG GGGAGATGCT GCGGAACAAG CTTGTCCGCC TCATGACACA	1200
55	CCTGGACACA GATGTGAAGA GGGTGGCTGC CGAGTTCTTG TTTGTCCTGT GCTCTGAGAG	126
	TGTGCCCCGA TTCATCAAGT ACACAGGCTA TGGGAATGCT GCTGGCCTTC TGGCTGCCAG	132
	GOGCCTCATG GCAGGAGGCG GCCCGAGGGC AGTACTCAGA GGATGAGGAC ACAGACACAG	138
60		

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•	ATGAGTACAA	GGAAGCCAAA	GCCAGCATAA	ACCCTGTGAC	CCCCACCCTC	GAGGAGAAGC	1440
	CGCCTAACCC	TATOGAGGGC	ATGACAGAGG	AGCAGAAGGA	GCACGAGGCC	ATGAAGCTGG	1500
5	TGACCATGTT	TGACAAGCTC	TCCAGGAACA	GAGTCATCCA	GCCAATGGGG	ATGAGTCCCC	1560
	GGGTCATCT	TACGTCCCTG	CAGGATGCCA	TGTGCGAGAC	TATGGAGCAG	CAGCTCTCCT	1620
10	CGGACCCTGA	CTCGGACCCT	GACTGAGGAT	GGCAGCTCTT	CTGCTCCCCC	ATCAGGACTG	i680
10	GTGCTGCTTC	CAGAGACTTC	CTTGGGGTTG	CAACCTGGGG	AAGCCACATC	CCACTGGATC	1740
	CACACCCGCC	CCCACTTCTC	CATCTTAGAA	ACCCCTTCTC	TTGACTCCCG	TTCTGTTCAT	1800
15	GATTTGCCTC	TGGTCCAGTT	TCTCATCTCT	GGACTGCAAC	GGTCTTCTTG	TGCTAGAACT	1860
	CAGGCTCAGC	CTCGAATTCC	ACAGACGAAG	TACTTTCTTT	TGTCTGCGCC	AAGAGGAATG	1920
20	TGTTCAGAAG	CIGCIGCCIG	AGGGCAGGGC	CTACCTGGGC	ACACAGAAGA	GCATATGGGA	1980
20	GGGCAGGGGT	TTGGGTGTGG	GTGCACACAA	AGCAAGCACC	ATCTGGGATT	GGCACACTGG	2040
	CAGAGCMANT	GTKTTGGGGT	ATGTGCTGCA	CTTCCCAGGG	AGAAAACCTG	TCAGAACTTT	2100
25	CCATACGAGT	ATATCAGAAC	ACACCCTTCC	AAGGTATGTA	TECTETTE	TTCCTGTCCT	2160
	GTCTTCACTG	AGCGCAGGGC	TGGAGGCCTC	TTAGACATTC	TCCTTGGTCC	TCGTTCAGCT	2220
30	GCCCACTGTA	GTATCCACAG	TGCCCGAGTT	CTCGCTGGTT	TTGGCAATTA	AACCTCCTTC	2280
50	CTACTGGTTT	AGACTACACT	TACAACAAGG	AAAATGCCCC	TCGTGTGACC	ATAGATTGAG	2340
	ATTTATACCA	CATACCACAC	ATAGCCACAG	AAACATCATC	TTGAAATAAA	GAAGAGTTTT	2400
35	GGACAAAAAA	АААААААА	AAAAAAAAA	. AA			2432

40 (2) INFORMATION FOR SEQ ID NO: 49:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1742 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:

50	GTCCTGCAGG	AGCTGCACGC	GCCCGAGGTG	OGCANGAACA	AGGAGCAGCG	AGAAGAGATG	60
55	TCGGGCTAAG	GCCCCGGSAC	GRGSGCCCC	CATCCTGCGA	CGGAACACGT	TCGGGTTTTG	120
	GTTTTGTTTC	GTTCACCTCT	GTCTAGATGC	AACTTTTGTT	CCTCCTCCCC	CACCCCAGCC	180
	CCCAGCTTCA	TGCTTCTCTT	CCGCACTCAG	CCCCCTCCC	CTGTCCTCGT	GGTGAGTCGC	240
	TGACCACGGC	TTCCCCTGCA	GGAGCCGCCG	GGCGTGRAGA	CCCCCTCCCT	COGTGCAGAC	300
60	ACCAGGCCGG	COCCCCCTCC	GTCCCCCGGG	GGCCCTGTGA	GAGAGGTGGY	GGTGACCGTG	360

	GTAAACCCAG GGCGGTGGCG TGGGATCRCG GGTCCTTACG CTGGCCTGTC TGGTCAGCAC	420
5	GTGCAGGTCA GGGCAGGTCC TCTGAGCCGG CGCCCCTGGC CAGCAGGCGA GGCTACAGTA	480
3	CCTGCTGTCT TTCCAGGGGG AAGGGGCTCC CCATGAGGRA GGGGCGACGG GGGAGGGGGG	540
	TGATGGTGCC TGGGAAGCCT GCKTGTGCAN CCGGTGCTTG TTGAACTGGC AGGCGGGTGG	600
10	GTGGGGGCTG CAGCTTTCCT TAATGTGGTT GCACAGGGGT CCTCTRAGAC CACCTGGCGT	660
	GAGGTGGACA CCCTGGGCCT TCCTGGAAGC CTGCAGTTGG GGGCCTGCCC TGAGTCTGCT	720
15	GGGGAGTGGG CATTCTCTGC CAGGGACCCA TGAGCAGGCT GCATGGTCTA GAGGTTGTGG	780
15	GCAGCATGGA CAGTCCCCCA CTCAGAAGTG CAAGAGTTCC AAAGAGCCTC TGGCCCAGGC	840
	CCCTCCGTGG GACAGCCCCG CCGCCCCTCC CCACCAGGGC TITGCAGATG TCCTTGAAAG	900
20	ACCCACCCTA GAGCCCTTTG GAGTGCTGGC CCCTCTGTG CCCTCTGCCC TGGTGGAAGC	960
	GGCASCACAA GTCCTCCTCA GGGAGCCCCA AGGGGGATTT TKTGGGACCG CTGCCCACAG	1020
25	ATCCAGGIGT TGGAAGGGCA GCGGGTAAGG TTCCCAAGCC AGCCCCAACA CCCTTCCCAC	1080
	TIGGCACCCA GAGGGGGCTG TGGGTGGAGG CCTGACTCCA GGCCTCTCCT GCCCACACCC	1140
	TCTGGGCTGA GTTCCTTCTT TCCCTTGGAC GCCCAGTGCT GGCCTTGGAG GACGGTCAGC	1200
30	TOGAGGATGG COGTGCCGGA GCCTGTCTTT GTACCACTGC AGCATCCCCC ACTTCTCCAC	1260
	GGAAGCCCCA TCCCAAAGCT GCTGCCTGGC CCCTTGCTGT AAAGTGTGAA GGGGGCGGCT	1320
35	GAGTICTCTT AGGACCCAGA GCCAGGGCCC TCAACTICCA TCCTGCGGGA GGCCTTGGCC	1380
-	GGGCACTGCC AGTGTCTTCC AGAGCCACAC CCAGGGACCA CGGGAGGATC CTGACCCCTG	1440
	CAGGGCTCAG GGGTCAGCAG GGACCCACTG CCCCATCTCC CTCTCCCCAC CAAGACAGCC	1500
40	CCAGAAGGAG CAGCCAGCTG GGATGGGAAC CCAAGGCTGT CCACATCTGG CTTTTGTGGG	1560
٠.	ACTCAGAAAG GGAAGCAGAA CTGAGGGCTG GGATATTCCT CATGGTGGCA GCGCTCATAG	1620
45	CGAAAGCCTA CTGTAATATG CACCCATCTC ATCCACGTAG TAAAGTGAAC TTAAAAATTC	1680
.5	AATCAAATGA ACAATTAAAT AAACACCTGT GTGTTTAAGA AAAAAAAAAA	1740
	œ	1742

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(2) INFORMATION FOR SEQ ID NO: 50:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1487 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

•	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:	
	GCCACGAGCC TCCGCGAACT GTGGAGTCGG CGGAGGGCTG GAATCAGCGT GGGCTCCAGG	60
5	TOGCTGGCAG COGGGTGGCA GAACTCTTCC GAGGCTCCTT GGGAAGAAGC TACACCCGAG	120
	GGAGCCGGAT GGGCCTCGAA AACCTGGCCC GCTCTGGTTC TGTACCATTG CAAGGGGAAC	180
10	CGTAAACTGA GCTTTTCTAA CGTGGGTTTC TGCCAAGTAC TTTTCCAGCT GCCCCCTTCC	240
10	CCCCAGCACA CAGGAGAGCC TCTGTGTAGC CAGCGCTTGA CAGTCGTTAG GTAGGTTGTA	300
	CTGTGTAGGG AGGAGCTCAA GATCATGAAT GGTTGTCACA GGAGAAAGCG GTTGCATCTT	360
15	TOCAAAACTA TATACCIGCT GIGGITIGIG TITICTITIC TGCTGAGTAA TGAAGIIGTA	420
	AGITCACACT GGCACATTCT CAGGCCTGTG CAGATTATTT GCACTTTATT TCATAGGTGR	480
20	ATAAGTGCTT TTTAGCTTTC TTTGTATATT GAGTTGCTTT TGAATTGCTT CCCATATTTT	540
20	TATTTCATAC AAACTGAACA ATTGTGGCCC CTCTATTTTA TTTATAAAGG TTCAGTGTAT	600
	CTITGCCTGC CTACATCAAT CTGCAAGGGA GTTGCAGAAA GCCTCATGTT CATCGAGCCG	660
25	TGAGTCACAA CCAATTTCTA AGCTGTTATA ACAAAAAAGT GTTTGCTTTT TTTCACAAGT	720
	AACTITAAAA GTGTAGTITA GAAAGAAAC ATTTTCAATA AAAAGACACT ACATTAATCC	780
30	TOGATOCTTG CAAATCCTAA AATMTATTCC TCCTCTAGCG TTGCACAGCT CTGTGTTGTA	840
50	TACACAGACT AGCTTTAAAA TITGTCACAT ACCACTTTAC CTTTACTTTT ATGTATCATT	900
	CCCCCGACTT CCTTACTGCA GGIGTGGGCA AGAAAACTTT TCCTTTAACA CTTTTCAACA	960
35	GCGGGCATAA AATTCTGCAG CTGAGGTCTT GAAGAATGCA GATGGGTACA GTATGTGTTC	1020
	GAGCTCACAG TGTGTATTGA CTAACCTAGT TCCTTTTTTG CTTTTTTTGG TATTGTCTTG	1080
40	TTAAAAGTGA CTCCCAGGTA GCAACTCTCT TTTTTAAGGG TGGGAACGAA AGGGACGTAG	1140
	GAAGAATAGA TCTAGATTAT TTAACAGTCT TCGATAGAGT TTGAAAGCTT TCTTCTTCAT	1200
	TCAATTTTGG GCAAAATACT GCCTCTGCAT TTGTTCATAA CAAAAAGATT AGATTAATAA	1260
45	GTAGCTTTTG TIGGTGGAAA TTACCAGCTC TATAAGTCAC CCTTGGTGGT TCATGGACCT	1320
	CTGATTAGCT TGGGTTTTGC AGTCTCATTG CCACATGTAT ATGTGGAGCC AATGGCCTTT	1380
50	TGGTGCTCAG CTGTTTACGT CTGACTCCTT GACTTCTTTG GTACAGTGAT GGAGTCAGAT	1440
	CTCATTAAGT GTGATTCTCC ATGGATATAA CCAGCCCCAA AAAAANG	1487

- (2) INFORMATION FOR SEQ ID NO: 51:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1328 base pairs
- 60 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:

)	GGCACGAGCT CGTGCCGAAT TCGGCACGAG AGAAGATTTG AAGAAGCCAG ATCCAGCTTC	60
	CCTGCGGGCT GCTTCTTGTG GGGAAGGGAA AAAGAGGAAG GCCTGTAAGA ACTGCACCTG	120
10	TOGCCTTGCC GAAGAACTGG AAAAAGAGAA GTCAAGGGAA CAGATGAGCT CCCAACCCAA	180
	GTCAGCTTGT GGAAACTGCT ACCTGGGCGA TGCCTTCCGC TGTGCCAGCT GCCCCTACCT	240
15	TGGGATGCCA GCCTTCAAAC CTGGGGAAAA GGTGCTTCTG AGTGATAGCA ATCTTCATGA	300
13	TGCCTAGGAG GTTCCTGACA TGGGACCCAT CTGCTCCTCC AGCCAACTCC TGTCCCTCAC	360
	ATCCCACCAT GGTGGCTCCT CCCACCTCCT CTGGATTTGT TCACTCTGAG ATCTGTTTGC	420
20	AGAGTGGGTG CTTAGCAGAC AGAGTGAAGC TGGCTGGGGG GCACAGTGGT GTGTAGTGCT	480
	GCTGTGTATC AAAAGACCAA GGTATTATGG GACCTGGTTT CAGAATGGGA TGGGTTTCTT	540
25	CACCTCATGT TAAGAGAAGG GAGTGTGTCC TGAAGAAGCC CTTCTTCTGA TGTTAAAATG	600
23	CTGACCAGAA CGCTCTTGAG CCCAGGCATC GTTGAGCATT AACACTCTGT GACAGAGCTG	660
	CAGACCCCTG CCTTGAGTCT CATCTCAGCA ATGCTGCCAC CCTCTTGTCT TTCAGAGTTG	720
30	TTAGTTTACT CCATTCTTTG TGACACGAGT CAAGTGGCTC ACAACCTCCT CAGGGCACCA	780
	GAGGACTCAC TCACTGGTTG CTGTGATGAT ATCCAGTGTC CCTCTGCCCC CTTCCATCCC	840
35	CAACCACATT TGACTGTAGC ATTGCATCTG TGTCCTGTTG TCATTTATGT TAACCTTCAG	900
-	GTATTAAACT TGCTGCATAT CTTGACATAT CTTGAGATTC TGCATGTCTT GTAAAGAGAG	960
	GGGATGTGCA TTTGTGTGTG ATGTTGGATA GTCATCCACG CTCAGTTTGG ACCATTGGAG	1020
40	GAACTTAGTG TCACGCACAA ATGGGGCTAT TCCTACGCTT AGAATAGGGC TTGTCTGCCC	1080
٠.	ACTITAGAAG AGTCCCAGGT TGGTGAGCAT TTAGAGGGAA GCAGGGCAGA ACTCTGAACG	1140
45	ACAATACGTC TCTCTGAGCA GAGACCCCTT TGTTCTTGTT ATCCACCCAT ATGGACTTGG	1200
	AATCAATCTT GCCAAATATT TOGAGAGATT GTGTGGATTT AAGAGACCTG GATTTTTATA	1260
	TTTTACCAGT AAATAAAAGT TTTCATTGAT ATCTGTCCTT GAAAAAAAAA AAAAAAAAAA	1320
50	ስ እ አርጥርር እ	1328

(2) INFORMATION FOR SEQ ID NO: 52:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1856 base pairs

(B) TYPE: nucleic acid

60 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ □ NO: 52:

	(XI, DECEMBER 2000)	
5	GAATTCGGCA CGAGCTCTGC AACATTSCAA ATTAACTTTGC AGTCGAGGGT TCCGCTGCCC	60
	CCTAGATTAA ATTCCCCGGG CTGAAACTGA GTTGCAGATT TACAATATCA TATTTTAAAT	120
10	TOCTGTCTTC AATTAAACCA TTTATGALTA TAALTAATTI TCAGGATGTC GATGCATGCT	180
10	THTCCAGGCC THOCTTCTTT GTACAAAAST AAATGTCCAT AAASCGTTTC ACTTATATTC	240
	TTCAAACATG ATGCTAATTT AAATTAALTA CTCCTATGA TALTSTTATTA TTCCTATGAT	300
15	THYGCCACTG TTATTAGTTC TCTCAAAAAT ACATCTAGGG AAGAGGATTA TTTTAAGTRA	360
	TYTGATTATC TYTTATTTAT TYCTCATTA CITTAGAAAT TOGTTCCATT	420
20	GGTTGGCATT GATACAGTAA ATTTGTAAAT GAGGAGACAA TATAAAAAAT CTAAATTACT	480
20	TGTGCTTAAT GACTGTAGCA GAATSCOTTT TCTCTAAATT AGALTGTCTT TCTTGCAGTT	540
	TAGTITGATA GATTIGCAAG CTATGCTGCT TCCATGAAGT TAGTIGCGCT GGTAGGAACG	600
25	CAGGCTTCTT TGTCTCTGGT TGTAGCTTGC ATGATCGCCC CALTIAGGCAG ACAACGTAGC	660
	COGAGATCAC AAATCAGGCC CTTOGTGTAG TTGCTAGTGT GTGGAGGTGC AGAGAGGTTG	720
30	GCAGAAACTG ACCTCACTGG GCAAGGGTGG CCATGGACCT GATTCTTTAA TGCACTCTAT	780
50	GTGTTCAGGA AGCCACAGGC CATATTTGAC TCTGAGAAAA AAAACAAGAG GAAAAAACCCC	840
	ACAAAGTATA ACAACCOCTT AAGATALATO TATTYLAAAS TGAAATTAAT TTTTCAGTTT	900
35	ATACCATTGG CCAATTACAA GATAAAAATG TTCAATTTGT TTLAGAATCC TTTGTTGACT	960
	TOTOTTTCA TOTOTTGCTA TYPATATUTG TOACTGTTAG TOAACAAGT CTTATTTGCT	1020
40	GAGGAAGGAC TITECTSCAC TEACTGEACE ACATCAAACA CESSGGAGGG TGGTGTTTAA	1080
40	CTTTTTAAAA AATGTTATTC TGATTATAAD AATAATATTG GCTTTTTCA TGAAAAGAGC	1140
٠.	GCCACCTTGC AAGGTTTAGT GAGATTTATG GAAGTTGAAT ATTTAAGCAG GAATTGCTGC	1200
45	TAGCTCCAAA AATTTGCGAA GCAAAASCTA GCCCCAATTS SITTGGAAGT TTGAAACTGA	1260
	TTAACAGATT TOCATTTGAA GTGACTTCAG ACATTAGGTT CAGACATTAG TTAAAAATAG	1320
50	AAAGAGGAAT AAAGACATCT YTTCTCTCTA GAAAAGATAA CACCECAACT AATAATCCTT	1380
50	CCCACTITICA TIGAGATCAG CITGICIGAZ AACCIGATAZ GASIGIGAZA AIGATAAACA	1440
	TGATAATAGT GGTACTTTIG TAATTTIGCT GGTSCATTIA AGAAGATAGT AAAKGATGAG	1500
55	TTCAYCTTTT CTYCGAACAT YCCTATYCCT AGATGTAGTT TACCTCAAAT TGGGAATTAT	1560
	AACTGTCCTA ATTTTTGTTG TGTACCTTGA TGCCCCTTTT GCTTTAATAC CCACAGTGTA	1620
60	ACAATTAAAT ATCACACTAT GACATAIGAI TIAAGIAGGA TATUITAAAG ATAAATTTTA	1680

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•	GCCGTAAATG	TTTACTTCAA	AATGACTCCA	TATTTCAAAT	ATCTGTTTAG	ACTGTGAAGG	1740
	CCAAATAATT	TITAAGAAAA	CATTTGAAGA	GTAGTGTGTT	TGCATTTGTG	AATAATCTTA	1800
5	CTCACAGCAA	GTAAACGTAA	TAAAAGCCAA	CATTTAAGCC	АААААААА	ААААА	1856

10 (2) INFORMATION FOR SEQ ID NO: 53:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1558 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:

20	TGGGTATCCA	TTCCTGNAAT	TACTTTACTT	AGGATAATGG	CCTCCAGCTC	CGTCCAAGTT	60
	GCTGCAAAAG	GTATTATTTC	GTTCCTTTTT	GTGGCTGAGT	AGTATTCCAT	ogtgtatata	120
25	TACCACATTT	TCTTTATCCA	CTCATTGCTT	GATGGGCAGT	TAGGTTGGTT	CCACATCTTT	180
23	GCAATTGTGA	GTTGTGCTGC	TCCAGATATC	ATCTTTAACT	CCTTTGCCTŢ	CTCCACATAC	240
	ATTTCCAAGT	CCTGTTCATT	CTACCTCCAA	AATGTATCTT	GTATCCATTC	ATCTCTCTCC	300
30	ATCTTCAATC	TATTTCAATG	CCCCATCATC	TCTTGCATGG	AGGAGTGTAA	TAATTGGCTA	360
	ACTGGCCTGT	TCTTACATTT	талаатсала	AGATGTGACA	GGTGAAATGC	CTATTTCAGT	420
35	GTCCATTGAT	GGTTCTGCTT	ACACACCACC	TGGCTGCCTG	GTGTCGCAGT	GGCAGAGTTG	480
<i>JJ</i>	AGCAGTGTGA	AAAAGACTGC	TTGGCCCTTT	ACAGGGAAAG	CAGGTCCACT	CTCCCCTCTC	540
	AGGACGAGAG	CTCTGGGCAG	GCTCGGACAC	TGGCAGACCC	TGGTCCTGGC	TGGCCAAGGC	600
40	AGCAGGGTAT	GTGTTTCGGG	TCACTCACAG	GGCTCAGCAC	CACTCCTCAT	GGCTTCCTTA	660
	CTGTTTCGGC	AGAGGCTGAC	CCGCGGCTGA	TTGAGTCCCT	CTCCCAGATG	CTGTCCATGG	720
45	GCTTCTCTGA	TGAAGGCGGC	TGGCTCACCA	GGCTCCTGCA	GACCAAGAAC	TATGACATCG	780
43	GAGCGGCTCT	GGACACCATC	CAGTATTCAA	AGCATCCCCC	GCCGTTGTGA	CCACTTTTCC	840
	CCACCTCTTC	TGCGTGCCCC	TCTTCTGTCT	CATAGITGTG	TTAAGCTTGC	GTAGAATTGC	900
50	AGGTCTCTGT	ACCCCCACT	TTCTCTGCCT	TCTTCCAGGA	TCAGGGGTTA	GGGTGCAAGA	960
	AGCCATTTAG	GGCAGCAAAA	CAAGTGACAT	GAAGGGAGGG	TCCCTGTGTG	TGTGTGTGCT	1020
F.F	GATGTTTCCT	GGGTGCCCTG	GCTCCTTGCA	GCAGGGCTGG	GCCTGCGAGA	CCCAAGGCTC	1080
55	ACTGCAGCGC	CCTCCTGACC	CCTCCCTGCA	GGGCTACGT	TAGCAGCCCA	GCACATAGCT	1140
	TGCCTAATGG	CTTTCACTTT	CTCTTTTGTT	TTAAATGACT	CATAGGTCCC	TGACATTTAG	1200
60	TTGATTATTT	TCTGCTACAG	ACCTGGTACA	CTCTGATTTT	AGATAAAGTA	AGCCTAGGTG	1260

314

	TTGTCAGCAG	GCAGGCTGGG	GAGGCCAGTG	TIGIGGGCTT	CCTGCTGGGA	CTGAGAAGGC	1320
5	TCACGAAGGG	CATCCGCAAT	GTTGGTTTCA	CTGAGAGCTG	CCTCCTGGTC	TCTTCACCAC	1380
	TGTAGTTCTC	TCATTTCCAA	ACCATCAGCT	GCTTTTAAAA	TAAGATCTCT	TTGTAGCCAT	1440
	CCTGTTAAAT	TTGTAAACAA	TCTAATTAAA	TOGCATCAGC	ACTITAACCA	АААААААА	1500
10	ААААААААА	AAANAAAAA	AAAAGGGGGC	CGCTCTAGAG	GTCCAAGTTA	NGACGNGG	1558

15 (2) INFORMATION FOR SEQ ID NO: 54:

20

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 948 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:

25	талалатсат	GCTCTGTACC	ATCCTCACCG	TAGTCATCAT	CATCGCCGCG	CAGACCACGA	60
	GAACTACTGG	GATCCCTAAA	AACGCCCCTG	GTCCGGCCCC	ACTCTGCGCC	CCTCGATCTC	120
30	CCAGGCTCTT	TCTGCAGWCA	TACCGCGGAC	CCAATGGGCG	CCCTGCACAC	CCCTTTCTCG	180
50	GGCCGTCAGA	CTTGGATACA	TCGTAAACTC	CGCCTCCACG	GAACGTCTCG	CCTKGCGAGC	240
	AAGMTCGGAA	TCCAGTTCCT	CAGGAACCCC	TCCAAAACCC	ACACCCCCAG	GGACGCCGCT	300
35	TTCCGGGATC	CCGGSCAAAC	GCCGGACCCT	CAGTCGCTCC	AGGCCCCCTC	ACCCTCAAAG	360
	TGTAGCGCCC	CCAACCGAGC	AACCTCGGTT	TGGTCCCTAA	AACCCCGCCT	CCTCTATAAG	420
40	CACCGCCCCA	GCTCTGACAA	AACCCCGCCT	CCAGGTCGGC	AGGCTCCGCT	TCTTTTCTTC	480
40	TCCGCGGGGT	GATTCAGTCC	AGTGATTGGG	TTTGTGGCTC	CAGGCCTCGC	CCACAGACGG	540
٠.	ACAGACCCCT	CCCTTTCTTC	CGGCAAAAGG	ACCGAGCCCT	GGGTAGTAA	GGSCCCCACA	600
45	CTCCTGTTTT	TTGCAAGTAC	ATTTTTGTCC	YTCCTCCACC	CAGGTATCTG	CCTATTTTCT	660
	TGCTAATCCC	AGAACCTTTC	CTTTTGCTTT	TTTTAAGGAC	ATTTGGGAAG	TTCCTGGTGT	720
50	AGGACCCTTC	TCCCTGGGAT	AAGAAACCTG	CCTGTAAACG	CTCTGTAAAT	ACTCCCTTCC	780
30	ACCCATCCCA	GCCCCTGGGC	AGCCGGGCAG	AAGGGAATCC	AGGCTATGGA	CCTCCCAAGT	840
	CCCCCCTCCC	CGCTCCCCTC	ecceccccc	CCTTGTTCTG	ATCTGTGTGT	GAGTGTGTGT	900
55	GAACTTCTGA	AAGACAATAT	TAAAGAGACT	TAGTTGAAAA	AAAAAAA		948

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5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 990 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:	
10	GGGGAACTGC AGTGACAGCA GGAGTAAGAG TGGGAGGCAG GACAGAGCTG GGACACAGGT	60
	ATGGAGAGGG GGTTCAGCGA GCCTAGAGAG GGCAGACTAT CAGGGTGCCG GCGGTGAGAA	120
1.5	TCCAGGGAGA GGAGCGGAAA CAGAAGAGGG GCAGAAGACC GGGGCACTTG TGGGTTGCAG	180
15	AGCCCCTCAG CCATGTTGGG AGCCAAGCCA CACTGGCTAC CAGGTCCCCT ACACAGTCCC	240
	GOGCTGCCCT TGGTTCTGGT GCTTCTGGCC CTGGGGGCCCG GGTGGGCCCCA GGAGGGGTCA	300
20	GAGCCCGTCC TGCTGGAGGG GGAGTGCCTG GTGGTCTGTG AGCCTGGCCG AGCTGCTGCA	360
	GGGGGGCCCG GGGGAGCAGC CCTGGGAGAG GCACCCCCTG GGCGAGTGGC ATTTGYTGCG	420
25	GTCCGAAGCC ACCACCATGA GCCAGCAGGG GAAACCGGCA ATGGCACCAG TGGGGCCATC	480
25	TACTTCGACC AGGTCCTGGT GAACGAGGGC GGTGGCTTTG ACCGGGCCTC TGGCTCCTTC	540
	GTAGCCCCTG TCCGGGGTGT CTACAGCTTC CGGTTCCATG TGGTGAAGGT GTACAACCGC	600
30	CAAACTGTCC AGGTGAGCCT GATGCTGAAC ACGTGGCCTG TCATCTCAGC CTTTGCCAAT	660
	GATCCTGACG TGACCCGGGA GGCAGCCACC AGCTCTGTGC TACTGCCCTT GGACCCTGGG	720
25	GACCGAGTGT CTCTGCGCCT GCGTCGGGGG NAATCTACTG GGTGGTTGGA AATACTCAAG	780
35	TTTCTCTGGC TTCCTCATCT TCCCTCTCTG AAGGACCCAA GTCTTTCAAG CACAAGAATC	840
	CAGCCCCTGA CAACTTTCTT CTGCCCTCTC TTGCCCCANA AACAGCANAA GCAGGANANA	900
40	NACTOCCTOT GGCTCCTATC CCACCTCTTT GCATGGGAAC CTGTGCCAAA CACCCAAGTT	960
	TAAGAAAAA ATAAAACTGT GGCATCTCCA	990
45		
45	10) TITODIOTECH TOD OTO TO NO. 56.	
	(2) INFORMATION FOR SEQ ID NO: 56:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1603 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
**	(D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:	6
	GGTCGACCCA CGCGTCCGGC CCGCCGGCTC CGGAGCGGGCT CTGCCTTCCC GAGCGCGGGA	12
۷0	CCGCCCCCTG GGGGAGGAGG GCGAACGACG CGGCGATGGC TCCGCGGGCA CTCCCGGGGT	12

٠	CCGCCGTCCT AGCCGCTGCT GTCTTCGTGG GAGGCGCCGT GAGTTCGCCG CTGGTGGCTC	180
	CGGACAATGG GAGCAGCCGC ACATTGCACT CCAGAACAGA GACGACCCCG TCGCCCAGCA	240
5	ACGATACTOG GAATGGACAC CCAGAATATA TTGCATACGC GCTTGTCCCT GTGTTCTTTA	300
	TCATGGGTCT CTTTGGCGTC CTCATTINGC CAMCTINGCTT NAAGAAGAAA GGCTATCGTT	360
10	GTACAACAGA AGCAGAGCAA GATATCGAAG AAGAAAAAGG TTGAAAAGWT AGRATTGAAT	420
10	GACAGTGTGA ATGAAAACAG TGACACTGTT GGGCAAATCG TCCACTACAT CATGAAAAAT	480
	GAAGCGAATG CTGATGTYTT AAAGGCGATG GTAGCAGATA ACAGCCTGTA TGATCCTGAA	540
15	AGCCCCGTGA CCCCCAGCAC ACCAGGGAGC CCGCCAGTGA GTCCTGGGCT TTGTCACCAG	600
	GGGGGACGCC AGGGAAGCAC GTCTGTGGCC ATCATCTGCA TACGGTGGGC GGTGTWGTCG	660
20	AGAGGGATGT GTGTCATCGG TGTAGGCACA AGCGGTGGCA CTTTATAAAG CCCACTAACA	720
20	AGTCCAGAGA GAGCAGACCA CGGCGCCAAG GCGAGGTCAC GGTCCTTTCT GTTGGCAGAT	780
	TTAGAGTNAC AAAAGTGGAG CACAAGTCAA ACCAGAAGGA ACGGAGAAGC CTGATGTCTG	840
25	TTAGTGGGGC TGAAACCGTC AATGGGGAGG TGCCGGCAAC ACCTGTGAAG AGAGAACGCA	900
	GTGGCACAGA GTAGCAGGTG AGCCGTGGTT TTGGTGACAT TGGGGGCAGA GTGGTGCAGG	960
30	GTGAGGAGAA GGTACTTGGA GCCTCCCAGG TGCTGTGGCA GCATAGGAAT GGTATTTGAC	1020
	AGGGAAGTGG GAGAGCTTTC CTTGACCCAG GAAGACTGAG GGGGACTGAA CATGATTACT	1080
	TGTCTGCCTA GAGCTTCTTG TAAAGAAGTC ACAAACTTAG TGCCTCCAGG GGCTTGGCTG	1140
35	TOTGATAATG AGGATAGAGG ATTACTTGTG AGGCAATGTG GCATGGTGGG GATTGTGGCA	1200
	AACTAGAATT CACATCACCC ACCATATAGG GCTTGCATTA CCACGAGGCA GAAAGCACCT	1260
40	AGIGITGCIG CATCITCITA CGCAAAAAAG ACAAAATCCA GACTICTAAA AIGIAAAATC	1320
	ACTGATTTTC GATATTGGCA GCTTACTTTT TTTTTTTAAA CAACCATGCA GGCCAAATGA	1380
٠.	CTTGTAATCT TGTCACCATT TTTAGGTAAA CTGTGACTTG AAAAAGTCTG GAGCAAACAA	1440
45	ACCAATGCTT TITCCTTTTA TICTGTTGGR AACCAGTFTT CTTTGTGTCA CAGTTYTGAA	1500
	ACCTCAATAC GAATATTTCT CTTCCCACCA AATATTTTGA GGCAATTGAA AAGCCACAGT	1560
50	GATTTATTIC TIGATITICC AATTTTAATT TIGCAAGACA ATT	160

(2) INFORMATION FOR SEQ ID NO: 57:

55

60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1052 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:

5	TACAGCTCAG GATGCCTGTA ACATTGTCAT CTCTGGGCTT CTGGGTCCTG CTTAGCCTGC	6
	TTTTTCCCTG GAGGACTGAC CAGGGATGCG GCCCAGCAAC ATGTTACTAA ATCATACTCT	12
	CCTCCCTACC TTTCCCAGAC CTCTCACTCC TGCCTGGTGT TCCAACCCGT TCTGTGGCCA	18
10	GAGTATACAT TTTGGAACCT CTTCGAGGCC ATCCTGCAGT TCCAGATGAA CCATAGCGTG	24
	CTTCAGCAGN AAGGCCCGAG ACATGTATGC AGAGGAGCGG AAGAGGCAGC AGCTGGAGAG	30
15	GGACCAGGCT ACAGTGACAG AGCAGCTGCT GCGAGAGGGG CTCCAAGCCA GTGGGGACGC	36
IJ	CCAGCTCCGA AGGACACGCT TGCACAAACT CTCGGCCAGA CGGGAAGAGC GAGTCCAAGG	42
	CTTCCTGCAG GCCTTGGAAC TCAAGCGAGC TGACTGGCTG GCCCGTCTGG GCACTGCATC	48
20	AGCCTGAATG AGGCTGGCCA CCTGCCACTT TGCCCTGCCC	54
	MYCCTICCTT TICTIGGIGA AAGGCACCIC CITTCCIGAT AATGAATGGI GITCCCTITG	60
25	CTTGGCTGGG GAGCCCCCCA GGCCAGGTTT GCTGGCCATA GATACCTTTG GGCTGCCTGR	66
25	GACAGOCTCC TGAGGAGGAT TGAGGGTGAA AGTCTCCCAC GAGTACACTA AACCTAGGTC	72
	TGGTCACCAA TAGGGTTTGG AGAGCAAAGG GCCACAACTC ATCAGCTGCC TGTCTCTTAG	78
30	ATGCACTITC TITTICCACC AGCACATCCT TCAACACACA GAATITCAGG GAAGAGITCT	84
	CCCCAAAACC CTAGCTCTTT ACCCTTCCAT TTTAGCCTTC CACCCAGCTT CCACAAAAGA	90
35	TTTGGCTCTA CCTTGGATCT GCTAGTAAAT AACTAATAGG CAGGCAGTTA TTTGGGTAAG	96
33	GAAAAAAGGG GTGGGAGAGA CAGAAAATTT GCCCACTGCT GCTCCTCCCC TTGGSTYTCC	102
	ACCTGGGATT TGCTATTGAA TCTCTACCCT NN	105
40		
٠.	(2) INFORMATION FOR SEQ ID NO: 58:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 814 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:	
	ACINCANTOGO GGCCGCTCTA GAACTAGGGG ANCCCCCGGG CTGCAGGAAT TCGGCACGAG	6
55	CATAGACTIT TAAACTOGTA COGTTCTTAG AGATGGTCCT TGGCCTTCTG TTGTTGTTGT	12
	RGITTITITE TITITETICI TERECTICIE CTICITETE TETICICET CITICITET	18
60	TTTTTTTCA GAGTCTTGCT CTGTCACCAA GACTGGAGTG AAGTGATGTG ATCTCGGCTT	24

•	AAGGTCAGGT TAGGGCTCCT GTACCCATTC TGTTCCACCA CTGTTTGATC TCTCTGGCCT	900
5	CCCACCAGGA ATGCCGTTTC CTTTTTATGG ATCTGTTGGG AACCAGAGAG AATCAACAGA	960
	TCAATGACAT AGGATCCGAA GTGCAATGAT AGTCACTTCT AGTTTGGCAT TTCACAAACT	1020
	CTGNACAGCA AGGTATTGGT AGGTTACTCA ATTTCAAAAG GGCCCCATGG CCAAATATGT	1080
10	TTAGGAACCG CTGTTTGNAT TTCTTTTTTT GGAGACGCAT TGTATATAAT ATATGTCAAA	1140
	GGCTTTCGGA ATTCCTGCAG GAAAGAAATC AGCTTTGTTA AATCCNAAAA AAAAAAAAAA	1200
15	AAAAAAATAG ACTCG	1215
15		
	•	
20	(2) INFORMATION FOR SEQ ID NO: 60:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 478 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
25	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:	
30	ATTTCTTATG ACATGGGGGT TTGAATTGGT TGGCAAATGT TTAATTTTAA TATCCATAAT	60
20	CAGTGAGGTC CTGCTGGCTG TAATCATTAA TTGTGAAATC TAAGGAGCTT AGTTCATGGC	120
	TCTAGAATTT CACAGAAAAR TGYGMTATGA TACGAGCATT AAGTTTATTT CTTCTGATCT	180
35	TTGATGCAGC TTTGTTCAGT TTATCTGTTT TTGTATTTAT TGGTCATCTA CTTCCCATGC	240
	CAAAAGGGAC TGGTCTACAT AGCTGCGCTA AACACCTGAT CAAATCACTA AAAGAAAATG	300
40	TGTTACCTCT AATGAATTAT CCTGATTGTA AGTTAAAAAT CAATATTTCC CCGTAGTGAG	360
	GTTTGCTTTT TAAAAAGAAK KCTTAAAAAA AAAAAAAAAA AAACGAGTTN AAGAAAAGGA	420
٠.	AGCAAGCTCA GGTAAGGTGC ACACATTGGG CTAAGGAAGC TAGAGCCTGT GGAGANGC	478
45		
	(2) INTERPRESENTATION FOR STO ID NO. (1	
50	(2) INFORMATION FOR SEQ ID NO: 61:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 618 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
-	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:	
	TATGACCTTG ATAACCCCAA GTTNGAAATT AACCTTCANT AAAGGGAACA AAAGCTGGAG	60
60	TICGCGCGCT TGCAGTICGA CACTAGTGGA TCCCAAAGAA TICGCCACGA GTCATAATGA	120

•	GCTACTAGGI AAGCCTTCTG GGACTTTCAG ATATTTTGGG GAAGATTGAT TTTTGTTCTT	180
5	ACATGCTGTG GACCCTTGGC CATCAAATGG TATGGGGAAG CTCATCCGTC TGTCTGTGAT	240
	GGTCATGTCA GTCAGGCGTC TTTTTAGTAT TTACTGGGTG CTCAGTACTG TGCCAGATGC	300
	TGTCGGGAGC CGTGGTGGTA TGGAGGAGGA GTGCTCCAGA GGACTCTGCT GTGTGGCAGG	360
10	CCAGCATAAA CAAGCCAAGG GGAAAAGGCA GGCATGGAAT AAAGGGGGGAG AATACCAGTG	420
	TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG	480
15	CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG	540
	TAGGACCINCA AGGCTTCTTIN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC	. 600
	AGTTTTGGGA AGCAAGGG	618
20		
	(2) INFORMATION FOR SEQ ID NO: 62:	
25	(a) and an	
25	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 751 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:	
	TOGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG	60
35,	TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA	120
	ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCONNT GTAAAATTCC ACCCCTGGAC	180
40	CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	240
	CTACAAGGAG ACTACGATGC CTGCCTTGGT CACCCTTCTC CTGCTCTTTC CATTGCTCCC	300
٠.	TCTGATGGAA GCCAGTTGCC ATGTGATGAG GTGCCCTATG GAGAGGCCCA CGTGACAAGG	360
45	TATTGTAAAA AGCCTCTGAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC	420
	TGAGATGAAT CCTGCCAACC TGAGCTTGGA GACAGATTCT CTCCCTATCC TGCCTTGGGA	480
50	TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	540
	GGTAAACTGG ACAGAATCCT GACCCACAGA AACTGAGATA ATGTTTGTTA TTTTAAGCTG	600
•	CTCAGTTTGT TACAGAGCAA TAGATAACTA ACTCAAACAC CATAAAATTC TAATATTTTA	660
55	TTCTATCACA CAAACCAGGT AATACCAAGT AAATGCCATT ACTATACACA TATTTTTGTA	720
	ACACAATTAC ATGTGATTTT TTAAGAAGGC T	751

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	(2) TECHNATION FOR SEQ ID NO: 05:	
5	(i) SEQUENCE CHARCTERISTICS: (A) LENGTH: 780 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPCLOGY: linear	
0	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 63:	
	CINENCASTICA CINETCCCCGA TICCCGGGTC GACCCACGCG TCCGGGTTGG CAACTCCTGA	60
	GGCCTGCATG GGTGACTTCA CATTTTCCTA CCTCTCCTTC TAATCTCTTC TAGAGCACCT	120
15	GCTATCCCCA ACTICTAGAC CTGCTCCAAA CTAGTGACTA GGATAGAATT TGATCCCCTA	180
	ACTEACTOTE TOCOGROCTE ATTOCTOCTA ACAGENTIGE CTGTGCTCTC CTCTCAGGGG	240
20	CAGCATECTA ACGOGGOGAC GICCTAATCC AACTGGGAGA AGCCTCAGTG GTGGAATTCC	300
	AGGCACTGTG ACTGTCAAGC TGGCAAGGGC CAGGATTGGG GGAATGGAGC TGGGGCTTAG	360
0.5	CTGGGLGGTG GTCTGAAGCA GACAGGGAAT GGGAGAGGAG GATGGGAAGT AGACAGTGGC	420
25	TOGTATISCOT CTGAGGCTCC CTGGGGCCTG CTCAAGGTCC TCCTGCTCCT TGCTGTTTTC	480
	TGATGATTIG GGGGCTTGGG ASTCCCTTTG TCCTCATCTG AGACTGAAAT GTGGGGATCC	540
30	AGGATGGCCT TCCTTCCTCT TACCCTTCCT CCCTCAGCCT GCAACCTCTA TCCTGGAACC	600
	TGTCCTCCCT TTCTCCCCAA CTATGCATCT GTTGTCTGCT CCTCTGCAAA GGCCAGCCAG	660
25	CTTGGGAGCA GCAGAGAAT AAACAGCATT TCTGATGCCA AAAAAAAAAA	720
35	GCGGCCGAAA GCTTATTNCC CTTTAAGTAA GGGGTTAATT TTTAGCTTGG GCACTNGGCC	780
40	(2) ENFORMATION FOR SEQ ID NO: 64: (i) SEQUENCE CHARACTERISTICS:	
45	(A) LENGTH: 588 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:	
	TTCCGAATTA ATCGACTCAC TATAGGAAWT GCCGTCGCCA TGACCCGCGG TAACCAGCGT	60
55	GAGCTCGCCC GCCAGAAGAA TATGAAAAAG CAGAGCGACT CGGTTAAGGG AAAGCGCCGA	120
		180
	CAGAAAAAGG CAAACGAGAA GAAGGAGGAA CCCAAGTAGC TTTGTGGCTT CGTGTCCAAC	24
60	CCTCTTGCCC TTCGCCTGTG TGCCTGGAGC CAGTCCCACC ACGCTCGCGT TTCCTCCTGT	30

	AGTGCTCACA GGTCCCAGCA CCGATGGCAT TCCCTTTGCC CTGAGTCTGC AGCGGGTCCC	360
5	TITITGTGCTT CCTTCCCCTC AGGTAGCCTC TCTCCCCCTG GGCCACTCCC GGGGGTGAGG	420
	GGGTTACCCC TTCCCAGTGT TTTTTATTCC TGTGGGGCTC ACCCCAAAGT ATTAAAAGTA	480
	ССТТІСТВАТ ТССАВАВАВА ВАВАВАВАВА ВАВАВАВАВА ВАВАВАВА	540
	AAAAAAAAAA AAAAAAAAA AAAANNCGGG GGGGGGCCCC CCCCCCCC	. 588
10		
	TO THE STATE OF TH	
15	(2) INFORMATION FOR SEQ ID NO: 65:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 774 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:	
25	TTTAAAGATG AAGAAATGAC AAGGGAGGGA GATGAGATGG AAAGGTGTTT GGAAGAGATA	60
25	AGGGGTCTRA GAAAGAAATT TAGGGCTCTG CATTCTAACC ATAGGCATTC TCGGGACCGT	120
	CCTTATCCCA TITAATTAAT TTCTCTGACA ATTCAATTAT TTTCTGTTAT TAATGTTGCC	180
30	ACTECTITCT GTTTGTCTGC ACTTTCTTGA TAAATATTTG CTATCGTTTT ACTCCAGTCA	240
	TTCGATGTTG CTGAGATTTA CATATGACTC TTGTCAACAT CTCATCTTTT GACCCAATCT	300
35	TATTCATTTA ATAAGAGGTC TCATTCATTT GCATGGAAAA ATGCTCATTG TATATTGCAA	360
33	AGTGAAAATA ACGAGTTGCA AAACAGTGTA TACATATATG TGTGTATATA TGTACACTTT	420
	ATTIGTACAT TICTATGIGA CATAATGCAA AGGAAAGIGT CIGATTITAT TATACACCAA	480
40	AGGITAACAG TGAATCTCTG TGTGATCTCT TTTTTTTTCT TTTTGCCTAT CTGCATCTTC	540
٠.	TCACTTGCCA AAAAATGAAT ATATGTTTAT GIGTGTATAT TACTTGTGTC ACAAAAAACC	60
45	CTAAAGTAGA CAGTAAAAGA ACTTGTCAAT CGCCTTTGGA AGGCAATGAA ACACTTAATA	66
43	AACTCTCAAT AACAGAAGCG TAAAAATGAA ATGTAAACCT CCAATTACCT CTGGATCTCT	72
	TAGCCAGAGT AATAAACTGG TAATTATTAC AGATAAAAAA AAAAAAAAAA	77
50		
	(2) INFORMATION FOR SEQ ID NO: 66:	
E		
55	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1866 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(C) STRANDELNESS: COUDIE	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:

•			
	ACCCACGCGT CCGGTCCTCT TCTTCAGCAC ATGCCAAAGC TGTTCCTCAC	GCCTGTGAG	60
5	ACAAGAGCAT CITGGATGTA GGACAATGGA AGAGTTAGAT GCCTTATTGG	AGGAACTGGA	120
	ACCCTCCACC CTTCAGGACA GTGATGAATA TTCCAACCCA GCTCCTCTTC	CCCTGGATCA	180
. 10	GCATTCCAGA AAGGAGACTA ACCTTGATGA GACTTCGGAG ATCCTTTCTA	TTCAGGATAA	. 240
10	CACAAGTCCC TTGCCGGCGC ANTCGTGTAT ACTACCAATA TCCAGGAGCT	CAATGTCTAC	300
	AGTGAAGCCC AAGAGCCAAA GGAATCACCA CCACCTTCTA AAACGTCAGC	AGCTGCTCAG	360
15	TYGGATGAGC TCATGGCTCA CCTGACTGAG ATGCAGGCCA AGGTTGCAGT	GAGAGCAGAT	420
	GCTGGCAAGA AGCACTTACC AGACAAGCAG GATCACAAGG CCTCCCTGGA	CTCAATGCTT	480
20	GGGGGTCTSG AGCAGGAATT GCAGGACCTT GGCATTGCCA CAGTGCCCAA	GGGCCATTGT	540
20	GCATCCTGCC AGAAACCGAT TGCTGGGAAG GTGATCCATG CTCTAGGGCA	ATCATGGCAT	600
	CCTGAGCATT TTGTCTGTAC TCATTGCAAA GAAGAGATTG GCTCCAGTCC	CTTCTTTGAG	660
25	CGGAGTGGCT TGGNCTACTG CCCCAACGAC TACCACCAAC TTTTTTCTCC	ACGCTGTGCT	720
	TACTGCGCTG CTCCCATCCT GGATAAAGTG CTGACAGCAA TGAACCAGAC	CTGGCACCCA	780
30	GAGCACTYCT TCTGCTCTCA CTGCGGAGAG GTGTTTGGTG CAGAAGGCTT	TCATGAGAAG	840
50	GACAAGAAGC CATATTOCCG AAAGGATTTC TTAGCCATGT TCTCACCCAA	CTCTCCTCCC	900
	TGCAATCGCC CAGTGTTGGA AAACTACCTT TCAGCCATGG ACACTGTCTG	GCACCCAGAG	960
35	TGCTTTGTTT GTGGGGACTG CTTCACCAGT TTTTCTACTG GCTCCTTCTT	TGAACTGGAT	1020
	GGACGTCCAT TCTGTGAGCT CCATTACCAT CACCGCCGGG GAACGCTCTG	CCATGGGTGT	1080
40	GGGCAGCCCA TCACTGGCCG TTGTATCAGT GCCATGGGGT ACAAGTTCCA	TCCTGAGCAC	1140
	TTTGTGTGT CTTTCTGCCT GACACAGTTG TCGAAGGGCA TTTTCAGGGA	GCAGAATGAC	1200
٠.	AAGACCTATT GTCAACCTTG CTTCAATAAG CTCTTCCCAC TGTAATGCCA	ACTGATCCAT	1260
45	AGCCTCTTCA GATTCCTTAT AAAATTTAAA CCAAGAGAGG AGAGGAAAGG	GTAAATTTTC	1320
	TGTTACTGAC CTTCTGCTTA ATAGTCTTAT AGAAAAAGGA AAGGTGATGA	GCAAATAAAG	1380
50	GAACTICTAG ACTITACATG ACTAGGCTGA TAATCTTATT TTTTAGGCTT	CTATACAGTT	1440
30	AATTCTATAA ATTCTCTTTC TCCCTCTCTT CTCCAATCAA GCACTTGGAG	TTAGATCTAG	1500
	GTCCTTCTAT CTCGTCCCTC TACAGATGTA TTTTCCACTT GCATAATTCA	TGCCAACACT	1560
55	GGTTTTCTTA GGTTTCTCCA TTTTCACCTC TAGTGATGGC CCTACTCATA	TCTTCTCTAA	1620
	TTTGGTCCTG ATACTTGTTT CTTTTCACGT TTTCCCATTT CCCTGTGGCT	CACTGTCTTA	1680
60	CAATCACTGC TGTGGAATCA TGATACCACT TTTAGCTCTT TGCATCTTCC	TTCAGTGTAT	1740

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5	ААААА						1866
	TAAATAAACT	GCCTTCTCCT	TTCAATAAAA	AAAAAAAA	АААААААА	АААААААА	1860
	TITIGITITI	CAAGAGGAAG	TAGATTTTAA	CTGGACAACT	TTGAGTACTG	ACATCATTGA	1800

10 (2) INFORMATION FOR SEQ ID NO: 67:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1152 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:

20	CTCAAGGATG	TAAAGGCTCT	GCAGATTTCG	GGAGGCCTGT	CTCCCAGCAC	CTGATGGGAC	60
	ACTITITICCC	CCACTGTAAA	TICIGGGIGT	ATCCTCCACT	GTATGCTGTC	ACCCCAAGGG	120
25	CAAGCACTGC	ATCTGCTTAG	TGAAGGATTT	ATTGTTCGGA	AGATACATTT	TCCCCTTKAG	180
23	CAGAGAGTGG	CGTATCCTGG	CAGTCTTCGG	TGAGCCAGTT	GTACCAGGAT	TATGAAATGC	240
	AGATGTTTAC	TGTGTCATTG	TTGCTGTCAT	TGCTACTGAG	GAGTACTGAC	CAGAATCATC	300
30	TGCAACTYTT	AGTTGGCAGA	GAGGACCACT	ATGGCGGGTA	GCTCTTTTCT	TTCCTGCCAT	360
	TGTGGGGATG	ATTCCAGGCC	AAAGATGATG	GARAAGTATG	GAAATCATCT	GAAAGGTTGA	420
35	AGCTTGGCAC	GTGAAGCCAT	TCATGACTTT	GTAAGGCAGT	TTTGCTGAAG	GCCAGTTCTG	480
33	CCCTGGGAGG	GACGGAGGTG	AATCCTCCTG	AGTACCTGTG	GTTTTCTTAC	TTCCTGCTGA	540
	ATTTACCTAA	GTGCCTGTTG	TTTGCTTGCT	GTGGAGGCTT	TCTGGTATTT	CATTICAGGT	600
40	GCAGATGCCT	TCACTTTCCC	ACCRAAAAAA	CCCCMACCAA	ACCTAAGACC	TTACTGCAAC	660
	TAAGTYTNCC	AAGTACTTTT	TAACCCAATG	GGATGAACAG	octotogict	GCTCAGATCA	720
45	CCCTGAGTGC	GTGTGAGAAG	GCMTNGGCTT	TGCCAGGAAA	TCCAGGAAGG	CAGGGCCGGG	780
45	CTGTGTTGGA	AGCTGGCTTA	GCTGGTGGG	CAGCCTTATT	TCAATTAAAA	GGGCATTGAC	840
	TGGGAGCAGC	AGTCCTGGAG	TTTGTTGCAT	TTCCTATTGC	CCTCAAAATG	AGAAACCAGG	900
50	AAAATAGCAG	ATTGGAGCCT	TCGAGAAGGC	: AGTAAATGGC	TGTTTTATT	GACAAAAGGA	960
	AAACATTTTA	CIGCCATCIC	ACTGATGGC	TCTCACTGAC	TTAAAATGAA	GCANGTTGT	1020
<i>-</i> -	AGTAAAAAAA	AAAGTCTACA	TTTTTCCACC	GCCACGTTCI	TATATCCTGI	TTGTCAGCCA	1080
55	CTGCTCANAA	GGGCATGTTG	TCTTGCGGAL	TANAGGCGCI	· creerreeci	CGITTICCCT	1140
	ATAGGTTGGG	TG					1152

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(2) INFORMATION FOR SEQ ID NO: 68:

5 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2483 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:

AGCAGGCGGT GCGCTGGGGG CGGGAGCAGC GCGKAGCCCG GCTCGGCCAC ACCGATCGCC

CGCCGCCATG GGCTCCTCGC AAAGCGTCGA GATCCCGGGC GGGGGCACCG AGGGCTACCA 15 CGTTCTGCGG GTACAAGAAA ATTCCCCAGG ACACAGACCT GGTTTGGAGC CTTTCTTTGA 180 TTTTATTGTT TCTATTAATG GTTCAAGATT AAATAAAGAC AATGACACTC TTAAGGATCT 240 20 GCTGAAASCA AACGTTGAAA AGCCTGTAAA GATGCTTATC TATAGCAGCA AAACATTGGA 300 ACTGCGAGAG ACCTCAGTCA CACCAAGTAA CCTGTGGGGC GGCCAGGGCT TATTGGGAGT 360 GAGCATTCGT TTCTGCAGCT TTGATGGGGC AAATGAAAAT GTTTGGCATG TGCTGGAGGT 25 GGAATCAAAT TCTCCTGCAG CACTGGCAGG TCTTAGACCA CACAGTGATT ATATAATTGG 480 AGCAGATACA GTCATGAATG AGTCTGAAGA TCTATTCAGC CTTATCGAAA CACATGAAGC 540 30 AAAACCATTG AAACTGTATG TGTACAACAC AGACACTGAT AACTGTCGAG AAGTGATTAT 600 TACACCAAAT TCTGCATGGG GTGGAGAAGG CAGCCTAGGA TGTGGCATTG GATATGGTTA 660 TTTGCATCGA ATACCTACAC GCCCATTTGA GGAAGGAAAG AAAATTTCTC TTCCAGGACA AATGGCTGGT ACACCTATTA CACGTCTTAA AGATGGGTTT ACAGAGGTCC AGCTGTCCTC 780 AGTTAATCCC CCGTCTTTGT CACCACCAGG AACTACAGGA ATTGAACAGA GTCTGACTGG 40 ACTITICIATI AGCICAACIC CACCAGCIGI CAGIAGIGII CICAGIACAG GIGIACCAAC 900 AGTACCGTTA TTGCCACCAC AAGTAAACCA GTCCCTCACT TCTGTGCCAC CAATGAATCC 960 AGCTACTACA TTACCAGGTC TGATGCCTTT ACCAGCAGGA CTGCCCAACC TCCCCAACCT 45 1020 CAACCTCAAC CTCCCAGCAC CACACATCAT GCCAGGGGTT GGCTTACCAG AACTTGTAAA CCCAGGTCTG CCACCTCTTC CTTCCATGCC TCCCCGAAAC TTACCTGGCA TTGCACCTCT 1140 50 CCCCCTGCCA TCCGAGTTCC TCCCGTCATT CCCCTTGGTT CCAGAGAGCT CTTCTGCAGC 1200 AAGCTCAGGA GAGCTGCTGT CTTCCCTCCC GCCCACCAGC AACGCACCCT CTGACCCTGC 1260 CACAACTACT GCAAAGGCAG ACGCTGCCTC CTCACTCACT GTGGATGTGA CGCCCCCCAC 1320 55 TGCCAAGGCC CCCACCACCG TTGAGGACAG AGTCGGCGAC TCCACCCCAG TCAGCGAGAA 1380 1440 GCCTGTTTCT GCGGCTGTGG ATGCCAATGC TTCTGAGTCA CCTTAACTTT GAACCATTCT 60

•	TTGGAATTGG	CGTGGTATAT	TTAACCACGG	GAGCGTGTCT	GGAAACGCAA	ACTATCATTA	1500
	ATTTCATACT	AGITTGTACC	GTATCTGTAG	CCATCCTGTA	AATAATTCCA	AGGGGAAAAC	1560
5	TAAACGAGGA	CGTGGGTTGT	ATCCTGCCAG	GTTGAGTGGG	GCTCACACGC	TAGGGTGAGA	1620
	TGTCAGAAAG	CGCTTGTATT	TTAAACAACC	AAAAAGAATT	GTAAGGGTGG	CTTGCTGCCA	1680
10	GGCTTGCACT	GCCGTTCCTG	GGGTGTGCA	TCTTCGGGAA	AGGTGGTGGC	GGGGCGTCCA	1740
10	CTAGGTTTCC	TGTCCCCTGC	TGCTCCTTCC	GTAAGAAAAT	GAAATATTCT	ATGCCTAATA	1800
	CTCACACGCA	ACATTTCTTG	TACTTTGTAA	GICGITIGCG	AGAATGCAGA	CCACCTCACT	1860
15	AAACTGTAAA	COGTAAAGAG	ATTTTTACTT	TTGGTCTCCG	TGAGTCGCAT	CTCTACTAAG	1920
	GTTTACACAG	GAATTCCACC	TGAAGACTTG	TGTTAAAGTT	CTACAGCGCG	CACTGTTAAC	1980
20	TGAACGICIT	TTTCTTCAGC	CTATACGCGG	ATCCTTGTTT	TGAGCTCTCA	GAATCACTCA	2040
20	GACAACATTT	TGTAACTGCT	GCTGTTGCTT	TCTACATACA	CCTTATAAAG	TGACATTTCA	2100
	AAAGAAATAA	GGTGCCACAG	TTTTAAACCA	GAAGGTGGCA	CTCTGTGGCT	CCTTGTAGTA	2160
25	TTATAGCTAT	ACTGGGAAAG	CATAGATACA	GCAATAAAGT	ACAGTAATTT	TACTTTTTTT	2220
	CTTGTGTTAC	ATCTAAATTA	CAACCCTTAA	TTGCCACGTG	TGCACTTACT	ACTCTCCAGT	2280
30	ATGTCTTATT	ACTOTOCAGT	ATGTCACGCA	TCTTTAACTT	TICACGICCT	ATGTTTGCTT	2340
30	TCTCCCATTT	TTAAGAGATG	GTAAGTTAAC	TGGAATTGAT	TTACTGAATG	AAATTAAATG	2400
	CAGATATCCC	TGTTTTTGAA	AAAAAAATA	AAAAAAAA	AAAAAAAAA	АААААААА	246
35	ааааааааа	ААААААААА	AAA				248

40 (2) INFORMATION FOR SEQ ID NO: 69:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 536 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:

50	GAGAAATGGA	GCTTTGTTAG	ATAAAAATTT	TTTCAACGCA	AACAGTCATT	TTCCAGTGAA	60
	AGGAGAGCGT	ATCCGCCGTA	GGATGGACTT	AGATCGTGTA	AAAGCTGAGG	CCACCGAGGA	120
55	TATAACCTCC	GGGGTCCTTT	GCCTCCTTTT	CCTTAGACTC	CCTCCAAACT	CCTCTATCTT	180
J J	TCCTTCAGCA	GTACTGGGCT	CCACGCGAAC	CTAGTCCTTT	GTCTTTACCC	TATTACCTTT	240
	CATAACATCC	TAGTTGAAAA	GTARTTATTC	AACCGCGTTT	GAAAATGAGA	ACAGGTTCAC	300
60	AGARGCTAGG	TTACTTGCGA	AGGTCGTTCA	ATTAGTAACC	AGTAACGCCA	GGACTGCCAG	360

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	TTTCTTGCTT CCGAATTCTC ATGGTAGCTT TCACCARGCT CCCCGTCMAA TGCTAACGTC	420
5	AACTACTGAA CTAGATTAGC AAAAAGGTCT TITAACAGAA TTCCTGGTTT TCAGAGAGAG	480
J	TTTCTTTCAT GAAGCGCCCC ATTTCTACAG AGGAAAATAA ACTCCAAGCA GCCAGT	536
10	(2) INFORMATION FOR SEQ ID NO: 70:	•
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 865 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:	
20	CCACGCGTCC GGCCTTTCTT GGCCAGAGGC GCCGGTTGGA CTCACGGGCG GGGCATGATG	60
	GGTAACAGGA CCGGTGGGGT CCCCAGGAAG TCCTAGAGGG GGTCGGGGTT TGGGTGGACA	120
25	AGCTTTCCTC GTCCTCTCCC GACAGAGCTG ACGTGTCCTG GGTTCCACCG GGAGCGGGCA	180
•	TTTCCACCGG ACGGGAGGGT TCGGGGGTGTC CGGGGCTGGG GAATACGTAG GGGTTGCCGC	240
20	GCGGTGTGGG GAGTTGGGGC GTGTGGCTGC AGTCCCGGGA GTTCTTGGAG GGGGTCGGCC	300
30	CACCGAGCTT CCGGACCGGC TGATCTGCCC GTAGCTTGCC GGANGGARGG CGGAGCTGAC	360
	TOTOCOTOCC TTOTOCCATO COCTOCAGTG GTGGGTACGG GCACCTCGCT GGCGCTCTCC	420
35	TCCCTCCTGT CCCTGCTGCT CTTTGCTGGG ATGCAGATGT ACAGCCGTCA GCTGGCCTCC	480
	ACCGAGIGGC TCACCATCCA GGGGGCCTG CTTGGTTCGG GTCTCTTCGT GTTCTCGCTC	540
40	ACTGCCTTCA ATAATCTGGA GAATCTTGTC TITGGCAAAG GATTCCAAGC AAAGATCTTC	600
40	CCTGAGATTC TCCTGTGCCT CCTGTTGCCT CTCTTTGCAT CTGGCCTCAT CCACCGAGTC	660
•	TGTGTCACCA CCTGCTTCAT CTTCTCCATG GTTGGTCTGT ACTACATCAA CAAGATCTCC	720
45	TOCACCOTGT ACCAGGCAGC AGCTCCAGTC CTCACACCAG CCAAGGTCAC AGGCAAGAGC	780
	AAGAAGAGAA ACTGACCCTG AATGTTCAAT AAAGTTGATT CTTTGTAAAA AAAAAAAAAA	840
50	AAAAAAAAA AAAAAAAAA AAAAA	865
55	(2) INFORMATION FOR SEQ ID NO: 71:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 932 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	
60	(D) TOPOLOGY: linear	

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	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:	
•	TCATCATATA CAAAGTTTTT CGTCACACTG CAGGGTTGAA ACCAGAAGTT AGTTGCTTTG	60
5	AGAACATAAG GTCTTGTGCA AGAGGAGCCC TCGCTCTTCT GTTCCTTCTC GGCACCACCT	120
	GGATCTTTGG GGTTCTCCAT GTTGTGCACG CATCAGTGGT TACAGCTTAC CTCTTCACAG	180
10	TCAGCAATGC TTTCCAGGGG ATGTTCATTT TTTTATTCCT GTGTGTTTTA TCTAGAAAGA	240
	TTCAAGAAGA ATATTACAGA TTGTTCAAAA ATGTCCCCTG TTGTTTTGGA TGTTTAAGGT	300
15	AAACATAGAG AATGGTGGAT AATTACAACT GCACAAAAAT AAAAATTCCA AGCTGTGGAT	360
	GACCAATGTA TAAAAATGAC TCATCAAATT ATCCAATTAT TAACTACTAG ACAAAAAGTA	420
	TTTTAAATCA GTTTTCTGT TTATGCTATA GGAACTGTAG ATAATAAGGT AAAATTATGT	480
20	ATCATATAGA TATACTATGT TTTTCTATGT GAAATAGTTC TGTCAAAAAT AGTATTGCAG	540
	ATATTTGGAA AGTAATTGGT TTCTCAGGAG TGATATCACT GCACCCAAGG AAAGATTTTC	600
25	TTTCTAACAC GAGAAGTATA TGAATGTCCT GAAGGAAACC ACTGGCTTGA TATTTCTGTG	660
	ACTOGTGTTG CCTTTGAAAC TAGTCCCCTA CCACCTCGGT AATGAGCTCC ATTACAGAAA	720
	GTGGAACATA AGAGAATGAA GGGGCAGAAT ATCAAACAGT GAAAAGGGAA TGATAAGATG	780
30	TATTTTGAAT GAACTGTTTT TTCTGTAGAC TAGCTGAGAA ATTGTTGACA TAAAATAAAG	840
	AATTGAAGAA ACACATTTTA CCATTTAAAA AAAAAAAAAA	900
35	CCAAATCGCC GCATAGTGAT CGTAAACAAT CT	932
40	(2) INFORMATION FOR SEQ ID NO: 72:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 996 base pairs	
٠.	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
45	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:	
50	COCCTGGCAC CATGAGGACG CCTGGGCCTC TGCCTGTGCT GCTGCTGCTC CTGGCGGGAG	60
J0	CCCCCGCCGC GCGGCCCACT CCCCCGACCT GCTACTCCCG CATGCGGGCC CTGAGCCAGG	120
	AGATCACCCG CGACTTCAAC CTCCTGCAGG TCTCGGAGCC CTCGGAGCCA TGTGTGAGAT	180
55	ACCTGCCCAG GCTGTACCTG GACATACACA ATTACTGTGT GCTGGACAAG CTGCGGGACT	240

TTGTGGCCTC GCCCCCGTGT TGGAAAGTGG CCCAGGTAGA TTCCTTGAAG GACAAAGCAC

GGAAGCTGTA CACCATCATG AACTCGTTCT GCAGGAGAGA TTTGGTATTC CTGTTGGATG

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•	ACTGCAATGC	CTTGGAATAC	CCAATCCCAG	TGACTACGGT	CCTGCCAGAT	CCTCAGCCCT	420
	AAGGGAACTG	AGACCAGAGA	AAGAACCCAA	GAGAACTAAA	GTTATGTCAG	CTACCCAGAC	480
5	TTAATGGGCC	AGAGCCATGA	CCCTCACAGG	TCTTGTGTTA	GTTGTATCTG	AAACTGTTAT	540
	GTATCTCTCT	ACCTTCTGGA	AAACAGGGCT	GGTATTCCTA	CCCNGGAACC	TCCTTTGAGC	600
10	ATAGAGTTAG	CAACCATGCT	TCTCATTCCC	TTGACTCATG	TCTTGCCAGG	ATGGTTAGAT .	660
10	ACACAGCATG	TTGATTTGGT	CACCTAAAAA	GAAGAAAAGG	ACTAACAAGC	TTCACTITTA	720
	TGAACAACTA	TTTTGAGAAC	ATGCACAATA	GTATGTTTTT	ATTACTGGTT	TAATGGAGTA	780
15	ATGGTACTTT	TATTCTTTCT	TGATAGAAAC	CTGCTTACAT	TTAACCAAGC	TTCTATTATG	840
	CCTTTTTCTA	ACACAGACTT	TCTTCACTGT	CTTTCATTTA	AAAAGAAATT	AATGCTCTTA	900
20	AGATATATAT	TTTAYGTAGT	GCTGACAGGA	CCCACTCTTT	CATTGAAAGG	TGATGAAAAT	960
	CAAATAAAGA	ATCTCTTCAC	ATGARAAAA	AAAAAA			996

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(2) INFORMATION FOR SEQ ID NO: 73:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 785 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:

35 GGCACGAGGG GCTTTGCGTA CACAATAGCT GCTAGGAGTA CCCAAAGCCT GARTACARCC 60 TECTEGTETC ATGGCCACGT GTGAGCAGGC CAGCGTCAMA CGGCTCGCTG TGACCCGTCC 120 40 CGRAGACTGA AATGGGCCTG GGTCTTCTCC TKGTCCTGTG ATWAAAGTCC TCTCTTGAAA 180 GTGGAGAGCA AAGGCACACA GAGGTGCGCG CTCACAAGAA TTCCTCCCGG TGACTGGGTA 240 ATCAATGITA CIGCIGITIC CTTIGCAGGA AAGACCACAG CAAGATICIT TCATICGICT 300 45 CCTCCTAGCC TGGGGGACCA GGCTCGAACT GACCCTGGAC ATCAAAGGAG GGATTATGTG 360 GCTGCTAAAG CCATCGCCCC ACAGCCCTGT TCACRTCTTG GTGCTTCTCT TTCCCAGAGG 50 480 CTGGTCCCAG CCAGGCACAC ACAAAAGGCA GATTCTCGTA AACSCAGCCT CCCTCCCTGG AGGCTGCCTC CTGCCCTGGA TCTGGAGTGG AGCTGCTCTG AGATTTTGAG TTCTTCTGCA 540 GAGATGATTA AATATATCCA AGAGACATTG GAAAACCTGC TGAACATTTT ACATTGGTCT 600 55 GCTCAGCACA TOGCTGGATG CGGATATTTC TATAATTCCA GAAAGTCACA CAGCTCCTCT 660 GTATGAGACC AGTGGGGGCC ATTTAAAAGA ACAGGATGAG AATCTAAGAT ATATTATTAA 720 780 60

ААААА 785

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(2) INFORMATION FOR SEQ ID NO: 74:

		CHARACTERISTICS:
/ i i	SECHIENCE	CHARACTERISTICS:

10 (A) LENGTH: 1069 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

	TCCTCACCAT	TCCCCTAGGN	CAGGTCCCTG	CAGGTCCCAC	ACTTCTCCCA	GGTCCCTAAA	60
20	CTTGGGTCGG	TCCTTTCCCT	GGAGTAGCTG	GNTCCTCCAG	TCGAGGTCCC	TGTTCAGTCG	120
20	GTTCTTAGGC	TCCTGCACAT	GAAGGTGTGT	GCCTGTGGTG	TGTGGGCTGC	TCTAGGAGCA	180
	GATACAGGCT	GGTATAGAGG	ATGCAGAAAG	GTAGGGCAGT	atgittaagt	CCAGACTTGG	240
25	CACATGGCTA	GOGATACTGC	TCACTAGCTG	TGGAGGTCCT	CAGGAGTGGA	GAGAATGAGT	300
	AGGAGGGCAG	AAGCTTCCAT	TTTTGTCCTT	CCTAAGACCC	<i>TGTTATTTG</i> T	GTTATTTCCT	360
30	GCCTTTCCGA	GTCCTGCAGT	GGCTGCCCT	GTACCCTGAA	CCTCATGAGC	CTCTAAGGGA	420
30	AAGGAGGAAC	AATTAGGACG	TGGCAATGAG	ACCTGGCAGG	GCAGARTACA	AGCCCAGCAC	480
	CAGTGTCCCA	GCCTTACTGG	GTCCTTACCC	TOGGCCAAAC	AGGGAGGGCT	GATACCTCCŢ	540
35	TGCTCTTCCT	AGATGCCCAC	CTCCTACAAT	CTCAGCCCAC	AAGTCCTCTC	CACCCTAGGG	600
	GCCTTCCTGC	ATGGCAATAA	CTCATAATCT	GATTTGGAGG	TTTGCCCTTT	ACAGGGGCAG	660
40	ATTTTCTGCT	CAGTTCAACA	ATGAAATGAA	GAGGAACTCC	CTCTTTCTAC	AGCTCACTTC	720
40	TATCAGAGGC	CCAGGTGCCT	CAGAGCCACA	TTGAGTTGCT	TTTTCTGGGA	TGAGGAAGTA	780
• .	GGGTTAAACT	CCCCAGTTTC	CTGAGGGAGG	CTCCTGACAG	GTGCCCTTTG	TCAGACCCTA	840
45	CCACAGCCTG	GATAGGCAGC	CACATTGGTC	CTCGCCCTTG	CTCGGNACTC	CGTGGTGGTC	900
	CTGCCCTTCT	CCCTGCATGC	CTGTGGGTCT	GCTCTGGTGT	GTGAAGGTCG	GTGGGTTAAC	960
50	TGTGTGCCTA	CTGAACCTGG	CAAATAAACA	TCACCCTGCA	. AAGCCAAAAA	AAAAAAAA	1020
30	AAAAAAAA	AAAAAAAA	. AAAAAAAAA	AAAAAAAA	AAAAAAAA		1069

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(2) INFORMATION FOR SEQ ID NO: 75:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 831 base pairs

(B) TYPE: nucleic acid

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(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75: 5 GGACATTAGA TCACTGTGGA CCTAAAACAA ACAAACAACT ATAAGGAAAA TGGCATTAGA 60 AATGGTCTGG GGATCAGTTT ATCACTGCAG TTGTTACATC ACCCCATGGT CTAAAATACA 120 10 GAGCTITAGT CTGTCTCTGT TTCAGTTCAT TTTACAGGAG GTGAACATCA CACTTCCAGA AAACTCTGTC TGGTATGAAA GGTATAAATT TGATATTCCT GTCTTTCACT TGAATGGCCA 240 GTTTCTGATG ATGCATCGAG TAAACACCTC AAAACTTGAA AAACAGCTCC TGAAACTTGA 300 15 GCAGCAAAGT ACTGGARGCT GACTGATGCC CTCATGATTT TCCACCCTCT CTTCCCATAA 360 AGCATCTTCC TAAGGAAATG AMCATGGCCT GATACTCATT TTGTCACTTG TACAGAGCCC 420 20 TAAGGATGTT CTGAATTCAG TGGTGCCAAA TAAATGTTGA CATTCCCCTT TTGGTTGATG 480 GAAGTATCAG TGTGGGAACT GTTTGCTTAA TGGCATTTTA TAAAATAAKA AKAKCATATT 540 AGCAGGGAGG GAGATGATGG AGGGAGGGAG AAGTCCATTT GTCTTATTTA TCCTTTTTGT 600 25 ATTAATAGAG AAGCACTTCA CAGTCACTGG CAATGCCATT TATAGGAAGA AGGTTCTGCA 660 TTCCTGCTGC TCCCGGAGGG CTTAACTTTT TAATGAAAGA ATAAATGCTC TTCCACTCAG 720 30 TAGATAAAGT GAAATGTGAA TTGTTAATAA CTGTGCACGG TCAATAAAGC GATGTTTTAA 780 831 35 (2) INFORMATION FOR SEQ ID NO: 76: (i) SEQUENCE CHARACTERISTICS: 40 (A) LENGTH: 590 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear

45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:

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TATATATAGA CNGTTAATAG TCGTGANTGN TGTGNACGAA CATTAACGGA AGTAGCATGT 60
AGCCAGTCGA ATAACNTATA AGGACAAAGT GGAGTCCACG CGTGCGGCCG TCTAGACTAG 120
TGGATCCCCC GGCTGCAGGA TTCGGCACGA GCTGCCAGGT GAGGAGCAGA GAGACTGTTC 180
CCTTGGGTGG AGAGGTGTGG GCATGAGAGC CACCCATTGC CAAGCAGCAA GAATGTTCGT 240
GCTTTTTTCC CTTCCAAAAT ATGCAGGGCT CAGGCTCCCA ATTCCGGGCC TGTCTGCTTT 300
GCTTGTGTTT CTCCTGTCCC TGTTCTCCCG GAGGGCCCAG GTGGAACTCA CGACAGGAG 360
GGAGACGCTT CCCAAAAACC TGCAGGGCTA TTTCCCAGAA TTTGGTTTTC AAGTACAAAA 420

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	CTTTTTGTCC	TGTAAGATAT	ATGCAGCCTC	ACAGAAGCAG	CCICIGCCIC	CACTTTACCA	480
	GCTACGTTTT	TATCTTAAGC	ACATGGGGCT	CCCTTAGAAC	TTACTCCACT	GATTTAAAAA	540
5	ааааааааа	AAACTCGAGG	GGGGGCCCGG	TACCCATTCG	CCCTAAAAGT		590

10 (2) INFORMATION FOR SEQ ID NO: 77:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1274 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

20	GAGCCACCAC	ACCTGGCCTG	GAAGGAACCT	CTTAAAATCA	GTTTACGTCT	TGTATTTTGT	60
	TCTGTGATGG	AGGACACTGG	AGAGAGTTGC	TATTCCAGTC	AATCATGTCG	AGTCACTGGA	120
25	CTCTGAAAAT	CCTATTCGTT	CCTTTATTTT	ATTTGAGTTT	AGAGTTCCCT	TCTGGGTTTG	180
23	TATTATGTCT	GGCAAATGAC	CTGGGTTATC	ACTITICCTC	CAGGGTTAGA	TCATAGATCT	240
	TGGAAACTCC	TTAGAGAGCA	TITIGCTCCT	ACCAAGGATC	AGATACTGGA	GCCCCACATA	300
30	ATAGATTTCA	TTTCACTCTA	GCCTACATAG	AGCTTTCTGT	TECTETETET	TGCCATGCAC	360
	TIGTGCGGTG	ATTACACACT	TGACAGTACC	AGGAGACAAA	TGACTTACAG	ATCCCCCGAC	420
35	ATGCCTCTTC	CCCTTGGCAA	GCTCAGTTGC	CCTGATAGTA	GCATGTTTCT	GTTTCTGATG	480
<i>JJ</i>	TACCTTTTTT	СТСТТСТТСТ	TTGCATCAGC	CAATTCCCAG	AATTTCCCCA	GGCAATTTGT	540
	AGAGGACCTT	TTTGGGGTCC	TATATGAGCC	ATGTCCTCAA	AGCTTTTAAA	CCTCCTTGCT	600
40	CTCCTACAAT	ATTCAGTACA	TGACCACTGT	CATCCTAGAA	GCCTTCTGAA	AAGAGGGGCA	660
	AGAGCCACTC	TGCGCCACAA	AGGTTGGGGT	CCATCTTCTC	TCCGAGGTTG	TGAAAGTTTT	720
45	CAAATTGTAC	TAATAGGSTG	GGGCCCTGAC	TTGGCTGTGG	GCTTTGGGAG	GGGTAAGCTG	780
73	CTTTCTAGAT	CTCTCCCAGT	GAGGCATGGA	GCTCTTTCTG	AATTTTGTCT	ACCTCACAGG	840
	GATGTTGTGA	GGCTTGAAAA	GGTCAAAAAA	TGATGGCCCC	TTGAGCTCTT	TGTAAGAAAG	900
50	GTAGATGAAA	TATCGGATGT	AATCTGAAAA	AAAGATAAAA	TGTGACTTCC	CCTCCTCTGT	. 960
	GCAGCAGTCG	GGCTGGATGC	TCTGTGGCCT	TTCTTGGGTC	CTCATGCCAC	CCCACAGCTC	1020
55	CCAGGAACCT	TGAAGCCAAT	CTGGGGGACT	TTCAGATGTT	TGACAAAGAG	GTACCAGGCA	1080
JJ	AACTTCCTGC	TACACATGCC	CTGAATGAAT	TGCTAAATTT	CAAAGGAAAT	GGACCCTGCT	1140
	TTTAAGGATG	TACAAAAGTA	TGTCTGCATC	GATGTCTGTA	CTGTAAATTT	СТААТТТАЎС	1200
60	ACTGTACAAA	GAAAACCCCT	TGCTATTTAA	TTTTGTATTA	AAGGAAAATA	AAGTTTTGTT	1260

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960 1020

1080

1133

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1274 TGTTAAAAAA AAAA

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(2) INFORMATION FOR SEQ ID NO: 78:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1133 base pairs

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	AGGATTTTTC	CTTGTTCAAC	CAAAATCTGA	GCATTCTTTC	TATGTTGAAA	ACACTGAAAA	60
20	ACTAATTIWA	GTTAATGAAC	TAGAAAGAAT	ATTGATTTTW	AAGAAACAGA	AAAATACTAC	120
20	TTATTTTCCT	TCTCAAATAA	CCTTTCTTTC	AAAAACTTCT	GGCTGAAGTA	TAACATGCTG	180
	GTAGTTAACA	TAAATCTTGT	CTTTCTCTTG	TTCTTTATCT	TTCTTTGTTA	TTTAGATGCT	240
25	TGTATAAATG	TCTTTTGTTT	TTATTAAGTG	CCTAATTGAC	AGAGCTTAAT	TTGAAGAAGT	300
	GCCCTAATTT	ATTGACCACT	TAAGAATTGC	CTTTATTGGG	GTATTTTATT	TGTTCCTGCG	360
30	TCTTTTTGAT	GTTGTTCAGT	CTACTCATCC	CTGTGAGTAT	GTGTGGGGGA	CAGCTGATAG	420
30	AAGGGAGGAG	AGTGTGTCTA	TGCTCAGGAT	TGCCCTTTAG	CCACTCAGCC	AGAGATCCAC	480
	AGGGAGCAAC	AAGGACAGTT	TCACATGCTT	AGACTTTCTT	GGAAGAAACA	GTGAGGAGGA	540
35	GTAAGTCGTG	AGTAGTGTCA	AGCTGGATGT	AGAATTGTCC	TAAGGCAGTT	GACCCCACCT	600
	TCCAACATGT	TTTCACTTTA	TTTGCCCCTC	CCTACATTTG	GGTTAGGTTC	CATTTGGATT	660
40	TGCAGCAATA	ATGACTTTAT	TTCTCTCTTG	GTCAGGATTT	GGCACATAAA	ATCCTTTTAT	720
40	TATAGAACTA	GCTATTTTAG	TTACATAGTA	ATGTAACTAA	TGGAGAGATT	TATAGAGAAT	780
	TTTGKTTTTG	CTGTCATATA	TGTCCATTTT	GGAGACAGAT	ATGATAGAAC	TAGAAATTAA	840
45	GTTGCATTTC	TGCAAGTGCC	ATTIGAATGA	ACTTCAAGTA	TCTTCTTAAT	TATTAAATTT	900

TCTGATGAAG GCATTGTAAC AAATATATAG TATTATTAAA TCTAATTAAT ATTTGGAAAT

ATTAATAAAT AGGTATTTTA TTTACTGTAA AAAGTCAAAC TTCATTATGT AGATAAATCT

TATTCTTTC ATTCTTCCC CTGTTTACAT CCTTTTTACA AAGCTTAGTC ACCAATTAAA

GCTTTCCTAT CAAAAAAAA AAAAAAAAA ACTCGAGACT AGTTCTCTCT CCT

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60 (i) SEQUENCE CHARACTERISTICS:

⁽²⁾ INFORMATION FOR SEQ ID NO: 79:

420

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	(A) LENGTH: 661 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:	
	GAATTCGGCA CGAGGGGAAA AGGATGCTGA ACGAGAGCAG AAAGCCTCTT TCCTTTGCTT	60
10	CACGCCTTTC CAGTCTTTAT TTTAAACTCG GGTTCCCTTT CTGTGGTCGC AGCAACCTTT	120
	ACTICACCTG CACTGCTGCT CCTGGGGGCT CCCCAGGCCT CCCTCTGCCT TTCTACCCAG	180
16	TESCTERACEG GATECCTETC TTESCTEGAC GCACCACTEC TCTCCTETCC CTCACCTTEG	240
15	CTTTTGCTGT GCCCTGCTCT GGGGTTGAAG CTGGCCCCATG TGTCCCCCGG AGTCATGGCT	300
	GCTCCTCCTG GGAGGCCTCT GTGTGCGTCA CGTCTTCCAC ACCTGGGGGC AGCTGGCGAG	360
20	CCCGTGCTCT GTTCCCCTCG GCTGCTTGGC ACAGAGYTGC AGCCTGGGAY TCTCCGTGGA	420
	CCCAGACTGG GGATTTTGCC AGGGGGGCGA TGGGAGGAGC AGGTGCTTTG CCTGGCGGCT	480
25	GTGTCTGCAT TTCTGGACGC CCCAGAGCAC AGAAGTTGCC GGCACTTTGA GGTCTTCCTC	540
23	GCCATGTCCC AGATTACATG AGTGACCGCT GCGAATATGT TTTCTTTTTT GTAATGGAGG	600
	CGTGTTTCAC ATATAGTAAA GCTCACCAAA AAGTAAAAAA AAAAAAAAA AAAAAACTCG	660
30	A	661
35	(2) INFORMATION FOR SEQ ID NO: 80:	
	(i) SEQUENCE CHARACTERISTICS:	
40	(A) LENGTH: 1378 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:	
45	ATTGGGTACC GGGCCCCCC TCGAAGTTTT TTTTTTTTT TTTTAATGAA AGCTCTCAAA	60
	TAAGCGATTT TATTCCTATC CATGATTGCA GACATTTACA AAACCATAAC ATCTGAGTTC	120
50	ACCTTAAAAA ATAACTTATA TAAAGCAGTG ATATACACAG CACAAAATAG TTCAGGGAGG	180
20	GGGCAGGAGC AACTIGIAAT AATTAAAATG TAAACGIGAA AAAAAGGATG GAATAAAAGT	24
	CCCTACTTAT TICTACTTAA GATGICATGI GATAATATTI TACAATGICC TGTGGGICAA	30
55	TGTATGTATG TGTATATGTC TGTATAACAT ACACATATAC AGTACATTCT CTTTCCCACA	36

CATATACATA CACACATAAT TATTTGCAGT TCAGTTTAGG GCAATTCTAA TATGCCACTC

CGTACAGTTG TTTGAATCAC ATTTGGACCC GCTTTCTTCA CAAAAGAGGG GAGAGAGCAG

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PCT/US98/11422

WO 98/54963

	GAAATAAAAA	CCTTCCTTTC	GTGTGACTGA	GATTCCTTTG	TTTAACTGTA	CACTGTGATG	540
	AATAATTTTC	TTCCGTAGTA	GTTCTGTGAA	GGGCTGACTC	ACTGTGGTTT	TCATGAGGAG	600
5	ACTTGGTAAT	GGATCACACG	CTCATTGTCA	TGCTAGGGGA	GTAACTCTCA	CTCTGAAAAG	660
	GATTTAAGAA	ATTTCCCCCC	ATTTCGCCAT	CATCCCTTGG	AGTGCCCGGT	TGATTACTCA	720
10	GGCTCATATT	ATTGGGAGAA	TICTTGGAAA	TACTGTCCAT	ATCTCCTGAG	CCTAAAGAGC	- 780
10	CATTCATGTG	ATGTGACTCC	ATTOCTCCTA	ATCCACCCAT	GGGACCATCT	GACCCAGGRC	840
	CCATTGGAAA	ATTAGGTCTG	TTAGGTCCAG	GAGGTACTGC	ATTCATTAAA	GTATACATGT	900
15	TATCACCAGA	GTTGGTTGAA	TCTGCTGGAC	TAGGCATGAT	CCCTCTTCCT	GGTGGCCCTC	960
	CACCTCCTGG	AGGACCTACA	TAATTCCCAG	GAGATGCTGA	GGAGTATGGT	ATTGAATTGG	1020
20	CATTTGTTGG	GTTTGGCCAA	GGTCTACCAC	CACCTGGACC	CATGTTCATT	CCAGGCATTC	1080
	CAGGGCCACC	TAAAGCATTC	AGTGGGGGTC	TCATTGCACC	TCCATAGTTC	TGTGGTCCTA	1140
	AGGGCACCAT	TCCTCTTGGA	GGAGTCATTC	TCTGCATTGG	CCCACCCATA	TTTGGATGTC	1200
25	CTTGTTGTCG	AGTTGGATCC	ATTCCACTGG	GGAGTAATGG	CTGACTTCCT	GGGACACCTC	1260
	CAAGTGCCTG	ATTAGGTATC	CTCAATGGGG	GCCTTGGACC	TCCAGGGTAC	CGAGGTGACA	1320
30	TAAAAGGGTA	ATCATGGAAG	GCTTTTGCTT	CACTTGAGTG	TTCACATGTT	TCACGTCT	1378

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(2) INFORMATION FOR SEQ ID NO: 81:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1440 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:

45	ACTTTGTCCA	AATGTGTCTG	TCACATGTAG	TCAGCTGNAG	AAATTTAAAA	TGAATTGCCA	60
43	AGTGAAGAGT	CTGTGGATTA	ATTGGCCGTT	AATTAACAGG	CTTTATCAAT	GTGTCCTCAA	120
	GGGAGAGGCC	CAACCCTAAT	TAAGGAGCTA	AACTTCCTGA	GTGAGGGGCT	GTGAGGATGG	180
50	AGGTGGAGGA	GGCATCTGGG	CCCCCTCCTC	GCCGGGCCAG	CAGATGGCGC	CTCCCTGGCT	240
	GAGCTGCCCG	CACCGCCAGT	TCCCTCATTT	CCACTCAGGA	AGGCAGAGAA	GGCAGAGTGA	300
-55	TCTCCTCAAG	GAAGAGCTTC	CCCAGCCTTC	GGGAGCAGCT	GGCAGGGCGT	CCGGGAATAA	360
.55	GCCCTACACG	CCCCCCCCIG	CCTCCAACTC	ACTAACCCTG	CCCCTCTTGT	CTTTCAGATT	420
	CAACGCGTTC	AACAGAAGCC	ATCCCCAGCC	CAGCTTAAAT	TATAAAGATA	GACAATAACT	480
60	CTGTTCCAAT	CTGCGTGGTG	CTTCTTTAGT	AAATACTGTA	CAGATTTTAC	CATGGAGAAC	540

336

	TTTTTTTTTA GTTTTTACCT TTTCTTAATT ACCCTTATTC CGAATGGACG AACACTTTCT	600
5	ACCACTGCTG ACCATTGTAA AATACCGTGT ATATAAATCC CATTGAAATA ATGCCCTGGA	660
J	ATAGAACATC TCAAATGCTG CTTAATTACA GACTCAGGTC GATTACTTGT ATTTCATGTA	720
•	ATGTTCCTCC AAGTTAGACA TCTGGTGCAA GACCAACCGG GAGACCATGG AATTGTCAAA	780
0	AGTACAAACT GACAGTGTGT ATATTTAATT TAAAGACTTA TITAAAAACT CACAAGCTCT	840
	CACCTAGACT TTGGAGAGCA GTCTGTTTTC TGTAATGTCT GATACTAGAA ACTAATTTGC	900
5	TTATTTTAGT TGTATTCAAG ATTTGAAGAT GTATTTTATA GACAAGTTCT GTTTTTGAAC	960
	TTTGTGGAAC TGTTCCAATC AATCAATTTC CCAGTTATGA TGAGTATTTA CATTATGAAT	1020
	GTATAACCCA GACATGATTT GTAAAGCCGA CAGTATGTTT CTATTACACA ACACTTTTTG	1080
20	ATACAGCGTC TCTTGTCTTC ACTGATACTG GAGTCTCCGT TGTCTGCNNG GTCCCTTCGA	1140
	GTTTCTAGTT ACAGACACAA TCATACTGIG ATTTTATTTT TAATATGGAT ATGCTATCAA	1200
25	ACTGTGATAC ACTTATAATT CACTGGTCCT GCATCAGGAG ATGGAGTGGG GAAAACTGTA	1260
	TTTAATACAG TTTGTATCTG AATAATCTGT ATGGTTTATA CAGTTTGTGT TGTTCAGAGA	1320
	TGTTTAAAGT TTGATCTTTG TTTTTCTAAA GATTAAAAAA GCACTTGCCC CACTGTAAAT	1380
30	ATACAGCATG TAAAATTTCT RTAGTATATA AATGGCAGCA AATCACAAAA AAAAAAAAAN	1440
35	(2) INFORMATION FOR SEQ ID NO: 82:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1381 base pairs (B) TYPE: nucleic acid	
40	(B) TYPE: MUCIEIC ACIG (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 82:	

60 45 CCCGGGCTGC AGGAATTCGK YACGAGGCCA GCAGTTGCTC CCAGTTCAGG AGGTGCTCCT GTACCCTGGC CACAGCCCAA TCCTGCCACT GCTGACATCT GGGGAGACTT TACCAAATCT 120 ACAGGATCAA CTTCCAGCCA GACCCAGCCA GGCACAGGCT GGGTCCAGTT CTGACCTGAG 180 50 CACGGTTTTT CCTCATGTGA CTTCTGGGAA GGCGCTCCCT CATCTGGGCC AAAGGAAGGA 240 300 GGACGAAGCC CTCCTCAGCT GGCCTGTGTT TGGGGCATGA ATCTCTCCTC TCCTCCTTGT 360 55 CTCGCTCTGT TGACAAACCG GGCATGTTTG GCAGTAAATT GGCACCGTGT CACACTGTTT CCTGGGATTC AAGTATGCAA CCAGAACACA GGAGAAGAAA AGCTCCAGGA TCCCTGTCCC 420 CATCTGTCCT CITGATGTGA GAGAGACTCT GAGACTTCTT CCATCGCAAT GACCTGTATT 480 60

337

	AAACACAAGC CCCCCAAGCA AAAGAAGAGG TIGAGTTTGC TGCCAGGATT CAGATCAGCC	540
	CTTCCCAGGG TCTGCAGGTG TCACATGATC ACAGTTCAGC GGGAGGCTTT CCGTACCCAC	600
5	ACTGGCTGTA GCACTTCAGT CCATCTGCCC TCCAGAGGAG GGTTTCTTCC TGATTTTTAG	660
	CAGGITTAGA GGCTGCAGCT TGAGCTACAA TCAGGAGGGA AATTGGAAGG ATTAGCAGCT	720
10	TTTAAAAATG TTTAAATATT TTGCTTTGCT AATGTGCTGA TCCGCACTAA CTCATCTTTG .	780
10	CAAAAGGAAC TOCTCCCTCG GCGTGCCCCA GCTGGGGCCT CTGAAGGGAT TCCTCACTGT	840
	GGGCAGCTGC CCTGAGCTTC AGGCAGCAGT GTTCATCTCT GGCCAGTTGT CTGGTTTCCA	900
15	TGTATTCTAG GCCAGGTAGG CAACACAGAG CCAAGGCGGG TGCTGGAAGC CAGACGGAAC	960
	AGTGTTGGGG CAGGAAGGTG GATGCTGTTG TCATGGAGCT GTGGGAGTTG GCACTCTGTC	1020
20	TGCTGGTGGC CCTCTCGGCT CACATGTTCA CAGTGCAGCT CCTGGCAGAC TTGGGTTTTC	1080
20	TCTTTGGTGG TTTCTAAAGT GCCTTATCTG CAAACAACTT CTTTTCTCCT TCAGGAACTG	1140
	TGAATGGCTA GAAGAAGGAG CTCAGTAAAC TAGAAGTCCA GGGTTGCTTG GTTTACTGGT	1200
25	TTATAAGAAA TCTGAAAGCA CCTCTGACAT TCCTTTTATT AACTCACCTC TCAGTTGAAA	1260
	GATTTCTTCT TTGAAAGGTC AAGACCGTGA ACTGAAAAAA GTGTTGGCCT TTTTGCGGGA	1320
	CCAGATTTTT AAGATAAAAT AAATATTTTT ACTTCTGTCA AAAAAAAAA AAAAAAATNT	1380
30		
30	С	1381
30	c .	1381
30 35	•	1381
	(2) INFORMATION FOR SEQ ID NO: 83:	1381
35	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs	1381
	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	1381
35	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	1381
35	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83:	
35	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT	60
35 40 45	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT AGCTCAGACT GGAGAGTAGC TTCAGGAAAA AAGACAAGTG GCCTAAGGAA ATCACGGCCC	60 120
35	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT	60 120 180
35 40 45	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT AGCTCAGACT GGAGAGTAGC TTCAGGAAAA AAGACAAGTG GCCTAAGGAA ATCACGGCCC	60 120 180 240
35 40 45	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT AGCTCAGACT GGAGAGTAGC TTCAGGAAAA AAGACAAGTG GCCTAAGGAA ATCACGGCCC CCAACTATCA TCTGAGGGCT AAAGATGAGA AGTAGATCAC TTAATAAGAC AAAAGCCTGT	60 120 180
35 40 45 50	(2) INFORMATION FOR SEQ ID NO: 83: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1706 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83: ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT AGCTCAGACT GGAGAGTAGC TTCAGGAAAA AAGACAAGTG GCCTAAGGAA ATCACGGCCC CCAACTATCA TCTGAGGGCT AAAGATGAGA AGTAGATCAC TTAATAAGAC AAAAGCCTGT AGGGGGAAAA GAAAGGATGT TTAAAAAGGAC AGAATGTTTC CCAAGGTAGA AATGACACTG	60 120 180 240

GGGGAAGAAT GAAGACACAG ACTCCTCTGT TCCCATTATC CCATCTAAGA CCCACACTCA

480

	CCTGGGGAAG CATCTGATTT AGAAATGTGG GTTAGTGTCC AGAGAATGGA AAAATAGACA	540
_	AGAGTCAAGG CTGGCAGGAT AACCTGTAAC AACAAAGGGT TTGAAAAATG AGGTTTGGGT	600
5	TAGGAGAGGG AGAGACAGAT AGCCAGAAAC ACACCAGTGA AGAGGAGAGA AAATGAGTAA	660
	AGGGAGAGCT AATTCCTTTT CCAGTGGAAA ATGAGTGATA TTCTGGACAT TCTTCAGAGG	720
10	CATCTACACG AAGTAGAAAT GTCACCGCTC CCTAATTTAC TCTACGTCTT CTAGAATCCC	780
	TCAATATTAT CCTTGGCTTC CAGGAAATCC AAGAAGACCC TGGAAGTAGA GTCCACCTTC	840
	TAAGAGAGA ATGTAAGAGG TGACCCCCAC CCACCTGATC TTCCTCGCTT TGTCCACTCC	900
15	ACGCACTGAG ACTTGACACA CCTAGTGGCC ACCTAGAACG TAGGTCCTTA AAATYTAGCC	960
	CCCCAGCCCC CAACCCATCT CTAGCCTGTC CACTCACCTG GTGAGGAACY TYTCCTGTGT	1020
20	CCACAGCYTT CTGCAGGAGT TGGCAACATG GCTCATAGAG CTCCCAGCGA GTCAGGTCAT	1080
	GAGTGCTTTG GGGGAGAAG GGGAATGTTA TACTGGAAAA GAACAGAGGG AACCAACTCC	1140
۰.	ACAGACACCA GTAAAAACGG GATGGGGAAG AGGAGGAAAG CCACTCACTT GTAGAAGGCA	1200
25	GAGAGGCGTT TCAGAGTGGC TGCCAGATTA TATACCTCAT CCTCATCTAG GAAGGACGAC	1260
	TGAGAAGGAA AGAAGATCCA CAATAGCATT TCCCCCAGAA CTCATCAGTC CACATCCCCC	1320
30	GTCTTGCAGC CCCTCCCACC CTTGTTTGGG GTGTCCCATT GTCCAGCCCC AGCTCCTACC	1380
	TGTAACAGCT CTTCAAGCTC CTGCTGGAAR CGGTCAGTCA GCAAATCTAC TAGCTGGCTG	1440
0.5	CGGGCAAAGT CCGCCCGGCT GAAGAAAGTG AATTCGGGAT TACAGAGCAG GTAAGAGCAT	1500
35	GCGCCCCAGC CTCAAGCACC GCTGGCTCTG CATGCTTCAC CACCACCTCC TGGAGTTGCT	1560
	GCAGGAACAG CTCCAGGTGC TGAGAAGAAA AGGCAGAAGA TGGTGTGCTG TGGGGATGGG	1620
40	AGGAGGACAC TCTTCTGGCG GGAAGTGGAA CGGGTTAAA AGCATTAAAC TTCAAGGATA	1680
	AGATGCCTAA RAAAAAAAA AAAAAA	1706
45		
	(2) INFORMATION FOR SEQ ID NO: 84:	

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 573 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:

GAATTCGGCA CGAGCTTGGT AGCCTTAGAA CTGCATGAGC TGCTTTACCA CTGGGAAACA 60
CGAGCACAGC CTAGCTTGAT TTTGTATGTG GTATCAGATC TAAGGTGGAT GGAATTCAGG 120

60

339

	ACTICCIGIC TACTCTTIGA TITTGTTTTA TITTTAGAAA TGTTTTATTT TGTTTTATTC	180
	ATTITATICAT CITCAGAGAC ATGGTCTGGC TCTGTTGCCC AGGATGGAGT GCATGGTGTG	240
5	ATCATAGGCC ACTGCAGTGT TGAGCTCCCG GGCTCAGGCG ATCCTCCTGC CTCAGCTYCC	300
	TTAGTAGCTG GGACTATAGG CACATGCCCT ACCATGCCTG GCTTTGTCTA CTTTTTGAAT	360
10	GATGTCYCAA ACTAGAAGGT CTATTAATTT AAAAAATTAA GGATAGCATG CCATAATTAA .	420
10	AAATAATAAC AGTGGGAAAA GGCACCTTCC AATGATTCAG ACATCAACTT GTGATTTAAA	480
	AAAACGAAAA ATAAATAATA GGAAAAAAAG GGGAAAAAGT TAAATAAA	540
15	AAAAAAAAA AAAAACTCGA GGGGGCCCG GTA	573
20	(2) INFORMATION FOR SEQ ID NO: 85:	
20	(2) INFORMATION FOR SEQ ID NO: 63:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 684 base pairs	
	(B) TYPE: nucleic acid	
25	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	• •	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 85:	
30	CTCTTTGGCT GTGTCTACCT CCTTCATCTG CTGCGCCGAC ATAAGCACCG CCCTGCCCCT	60
	AGGCTCCAGC CGTCCCGCAC CAGCCCCCAG GCACCGAGAG CACGAGCATG GGCACCAAGC	120
35	CAGGCCTCCC AGGCTGCTCT YCACGTCCCT TATGCCACTA TCAACACCAG CTGCYGCCCA	180
55	GCTACTTTGG ACACAGCTCA CCCCCATGGG GGGCCGTCCT GGTGGGCGTC ACTCCCCACC	240
	CACGCTGCAC ACCGGCCCCA GGGCCCTGCC GCCTGGGCCT CCACACCCAT CCCTGCACGT	300
40	GGCAGCTTTG TCTCTGTTGA GAATGGACTC TACGCTCAGG CAGGGGAGAR GCCTCCTCAC	360
	ACTGGTCCCG GCCTCACTCT TTTCCCTGAC CCTCGGGGGC CCAGGGCCAT GGAAGGACCC	420
45	TTAGGAGTTC GATGAGAGA ACCATGAGGC CACTGGGCTT TCCCCCTCCC AGGCCTCCTG	480
43	GGTGTCATCC CCTTACTTTA ATTCTTGGGC CTCCAATAAG TGTCCCATAG GTGTCTGGCC	540
	AGGCCCACCT GCTGCGGATG TGGTCTGTGT GCGTGTGTGG GCACAGGTGT GAGTGTGTGA	600
50	GTGACAGTTA CCCCATTTCA GTCATTTCCT GCTGCAACTA AGTCAGCAAC ACAGTTTCTC	660
	TGAAAAAAA AAAAAAAAAA AAAC	684

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(2) INFORMATION FOR SEQ ID NO: 86:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1036 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86: TOGAGGCAGA TOCACAGGAG AAAGGTTCCC GTCCGCACCC TCTCAGACCT GAGGCTGAGC TTGCAGTGAG GGCTTCTCCT CGGCCCCTCG CCCGCCCCCA GAGCTGCCAT CCCTGCTGTT -120 10 ACAAGCCAGA GGAGCCCGGA TGTGAGGCCC CAGATCACCT CCAGGGACTT GGGGTTCCCA TCTGAAATCC TTTATTTTTG TACCATGGGG TGGGCCCCGG GCTGAGAAGG AAGAAGCACC 240 15 CTCTCCCCGG CCTCCTCTGT CTGCACCCGT GGGGCTGTGA CTTACTCCTG CCTCCAGGGG 300 CGGGGCGGGG CCCCCTGGGA CCTCTTAAGG CCCAAGGTGG GCCCCAGGAC CTYTGGGCAG 360 AGTGGAYTGC TCATGGCAGA TGTGTGGCAA TGTCTGGCTG WGTCTTTCCG GCAMCTGCGT 420 20 YCCCTYTCCC GGGYTCCCCT GCTGCATGGT GGATGTGCTC CTTCCTGGCC CGGTCACATT 480 GCCTCCTTGA GCCTTAGTCC AGGGGGTCAC TYCTCCCACC CCACCTACCT CACAGGGTTG 540 25 TTGTGAGGGT GCACAGAGGA GCAAAGTCCC TGAAGGCCCT CAGGCAGTAT ATAGGGGCCG 600 CCCACCTTCA GCTGCCCTGG GATGGGAAGG ACCCAGCCCG ACCCCTGGGC ATAACACTGT 660 720 CTTTGCAAAT GGAGATTCAG GTATTGGGGA TGCAGGTTGT GGGGAGCTGG CCTGGCAGAG 30 TAGGGGTAGT TGGCTTGGCC TTCTCTTTGG TGATCCCACC CCCAGCCATT TGCATTGCTG 780 GCCCAGCGCC TGGCCTGGGG GGCGGGGAGA GGCAGCAGAA GGGGCTGGGC AGGGGCGGTG 840 35 GAGGACTCAG GAACTGCCCG GGGAGAGTGG GTATGGCGGC TGAGCCAGGG GCCCTCCTGT 900 GTTTGACTTC CCGGGATGGG TCCTTGCTTC TCAGCTGTGT CCGACCCCAC CATGTAATAA 960 1020 40 1036 CCCNGGGGGG GNCCCG

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(2) INFORMATION FOR SEQ ID NO: 87:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 908 base pairs(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 87:

TTAAACAAAT GGAATCATGC AATATGTGAC CTTTTGCGTC TGGCTTATTT TATTTAGCAT 60

AATGTTTTTG AGGTTCATCC AAGCTGTAGC ATGTATCAGC ACCTCATTTC TTTTTCTGGC 120

60 TGAATATTAT TCCATTATAT GGATTTACCA CAATTCATTT ACCTATTCAT CTTTTGTTTC 180

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	TOCTGTCTGG CTATTGTGAA TAATGCTTCG ATAAACATTC ATATACAAGT TTCTATGTGG	240
•	CTTTATGTTT TCATTTCTCT TGGCTATCTA CATGGGAGTA GAATTCTAGG TCATAATATA	300
5	ATTITATGTT TAACTICICA AAGAATTGCC AAAAGGTTTT TCATAGTGGC TGCATCATTT	360
	ACATTCCCAC CGGCAATGTA CAAGGATTTC TATTTTTCCA TATCCTTGCA CTTACCAACA	420
10	CTTCTTTTTK GIWATWATTT TGTTTTTCA TTATTGCCAC CCTAGTGGAT GTGAAATGGC	480
	ATCTTATTGT TITGATTTGC ATTTCTCTAA TGACAAATGA TATCATACTT TTTTTATGTG	540
1.5	CTTACGGATC AAAGGTATTT CCTTGGAGAA ATGTCCCTTC AAGTCCTTTG CCATTTCAAA	600
15	ATTIGGITAT TIGICTITTA TTATICAGIT TTAAGAAATT CIGGCCAGGC GCAGTGGCTC	660
	ACCTGTAATC MIAGCACTTT GGGAGGCCAA GGCGGGCAGA TCACTTGAGK TCAGGACTTC	720
20	GAGACCAGCC TGGCCAACAT GGTGAAACCC CATCTTACTA AAAATACAAA AATTAGCTGG	780
	GCGTGGTGGC AGGTGCATGT AATCNTATCT ACTCAGGAGG CTGAGGCAGG AGAATCGCTT	840
06	GAACCCAGGA GGCGGAGGCT GCAGTGAGCC AAGATCACGC CATTGCACTC TAGCCTGGGT	900
25	GACACAGA	908
30	(2) INFORMATION FOR SEQ ID NO: 88:	
	(i) SEQUENCE CHARACTERISTICS:	
35	(A) LENGTH: 655 base pairs (B) TYPE: nucleic acid	
<i>J J</i>	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:	
70	TGCACTGGTT CCTTCTCCCC AGCAAATACT GCCTTCTTGT TTTTCTCTGA TGTGGCAGGT	60
	GACTACAAAA TCCGCCTTGG TATTCTTCAA ATGCATATAT ATTCCTTTCT TGTCAGCTCC	120
45	CTCTCTTCCT AGATTAGAAA ACTGCCTCAT TTTCTGCTCA CTGGATGTGC AGTCCCAGCT	180
	TGTCTTCCTC TCCTCCCCCC CTGTTGCAGG TGTTCTTTTT TTTTTTCTTC TCTCCCCACT	240
50	GGGCAGCAAA AGTTGTTCCA CAGTGGAAAW TTAGGCATCC TCAAGTTTCY TCCCAGCTTC	300
3 0	TGCTGTGTTT TCTTAGAGTA AATTGCCAAT TTCTGTTTTT ACAGGAAATC CTTTTTTAAA	360
	AATGGAATCA GTGTGGTCCC CATCTACTCT GCAAAAATTG CATTTTTCTC TATTTTCAAA	420
55	TGAGATTTGT TCAAGTTTCA AAACCACGTG AAATAATAAA TGTATAGTAG TTTTCTTTTC	480

CTTGGGCATT GCTWGATATG TGAAATGGGT TTATGAAAAA TAATAAAATC ATAACGCTAT

TTGTTTGACT TTCAATTTCA TGGGAATTTT TCTCAGCTAA ACTCTAAATG GTGATTARGC

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ΔΔΔΔΔΔΔΔ	AAAAAAAACY	GRAGGGGGC	CCGGTACCAA	TICGCCCTAT	AATGA	655

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(2) INFORMATION FOR SEQ ID NO: 89:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1102 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 89:

15 TTTTTTTTT ACCATTAAA ATAAAATGAA AGTGACCTTC TGTTTATAAA AATCTTTGTC 60 TGCATCTCTG CTTATTTCCT TAGAAGAGAT TCCAAGAAGC GGTGAGTGAT TTCACGGCAG 120 CAGAGGGTTG GGACATATTA COGGCGGGA TCCCTCTTGG AGTGAGATGA CTCTCCGGAG 20 240 AGATTTAGTC GTCACCCTCG CGTGTGAGGC TGCGTCACAC CCCAGGGATG TGTCTATCAA GATGGAAGAT CTTTTACACG CTCTTGATTT TGTTTGSCTY TTTTTCTATT ACTAGTGAGA 300 25 360 AKGAAACTTT TTATATGATT ATTATCCATC ATAATCCAAC ACAAATTACT GCTTCATGTT CTTTTACTTT CCTGTGAAGG TTTTAGTGCC TTTTAAAAAT TGCTATATAT TAAGCTTGTT 420 AATACTTCCA TGCTGTATTT GTGGSCATCA RTTTCCCCGG GNACAGGCNT GCACATTTTG 30 CCTTCACACG CTGGGTGGTT TTTCATTTTC AMTTCTATTT CTCGTTCTTC TATCGTTTTA 540 TGTTCAGACG GGTTTCTCCG TGTAGAAAGC AGTTTATGAA GATTTACTTT CGACAGTCTT 600 35 CTCTCTACTT TCTACAGTGA ATTCTCTGAT GTGTCTGGGA GTTTGGGGGT CTGGGTAAGA 660 RICCICCICI CACCCIATIC ICIATIACGA ICCACAGCCI CATGCITIAI GARATIGGIG 720 780 GCCGGGARCG GGGGAGATTT GCGGATCCCC CAAGCCAGAC TTTATCCCCC TATCCCTGCC 40 TCTGGATCCC ACGTACAGGC CTGGGAACTC CCTGTGGGTA GGGGCCAATG GTCTCGCACT 840 CTCACCTGTA CCCCAGGGCT GGCACAGGAT GGTCAAGGAG AGAGGCTGCC CAAGCGCATC 900 45 CYTCTGGTGT CCCCCTGACA CGCCTCCAAA GTGAGCAGGT AGGTTTCAAC AGCCCCACGT 960 TOCAGGTOGG AGATGAAGCT CAGGGTGGAG ACCAGTATCT CACAGTTCTC TTTGCATGGC 1020 CGGGTACTTG TTAGTCAACT GATCAAGTGA AAATTCTAGC CCCAGAGGCA GGAGAATCCG 1080 50 1102 GAACAAAATT AAACCAGCCA GG

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(2) INFORMATION FOR SEQ ID NO: 90:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1533 base pairs

343

(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear

5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 90:	
	GGCACGAGCC GNCACGGGCA GCGCCCCATA GCGCCAGGGA CCCCCTGGCA GCGGGAGCCG	60
10	CGGGTCGAGG TTATGGATCC AGCGGGGGG CCCCGGGGGG TGCTCCCGCG GCCCTGCCGG .	120
10	TENCTGGTGC TGCTGAACCC GCGCGGCGGC AAGGGCAAGG CCTTGCAGCT CTTCCGGAGT	180
	CACGTGCAGC CCCTTTTGGC TGAGGCTGAA ATCTCCTTCA CGCTGATGCT CACTGAGCGG	240
15	CGGAACCACG CGCGGGARCT GGTGCGGTCG GAGGAGCTGG GCCGCTGGRA CGCTCTGGTG	300
	GTCATGTYTG GAGACGGCT GATGCACGAG GTGGTGAACG GGCTTCATGG AGCGGCCTGA	360
20	CTGGGAGACC GCCATCCAGA AGCCCCTGTG TAGCCTCCCA GCAGGCTCTG GCAACGCSCT	420
20	GGCAGCTTCC TTRAACCATT ATGCTGGCTA TRAGCAGGTC ACCAATGAAG ACCTCCTGAC	480
	CAACTGCACG CTATTGCTGT GCCGCCGGCT GCTGTCACCC ATGAACCTGC TGTCTCTGCA	540
25	CACGGCTTCG GGGCTGCGCC TCTTCTCTGT GCTCAGCCTG GCCTGGGGCT TCATTGCTGA	600
	TGTGGACCTA GAGAGTGAGA AGTATCGGCG TCTGGGGGAG ATGCGCTTCA CTCTGGGCAC	660
30	CTTCCTGCGT CTGGCAGCCC TGCGCACCTA CCGCGGCCGA CTGGCCTACC TCCCTGTAGG	720
50	AAGAGTGGGT TCCAAGACAC CTGCCTCCCC CGTTGTGGTC CAGCAGGGCC CGGTAGATGC	780
	ACACCTTGTG CCACTGGAGG AGCCAGTGCC CTCTCACTGG ACAGTGGTGC CCGACGAGGA	840
35	CTTTGTGCTA GTCCTGGCAC TGCTGCACTC GCACCTGGGC AGTGAGATGT TTGCTGCACC	900
	CATGGGCCGC TGTGCAGCTG GCGTCATGCA TCTGTTCTAC GTGCGGGCGG GAGTGTCTCG	960
40	TGCCATGCTG CTGCGCCTCT TCCTGGCCAT GGAGAAGGGC AGGCATATGG AGTATGAATG	1020
40	CCCCTACTTG GTATATGTGC CCGTGGTCGC CTTCCGCTTG GAGCCCAAGG ATGGGAAAGG	1080
	TGTGTTTGCA GTGGATGGGG AATTGATGGT TAGCGAGGCC GTGCAGGGCC AGGTGCACCC	1140
45	AAACTACTTC TGGATGGTCA GCGGTTGCGT GGAGCCCCCG CCCAGCTGGA AGCCCCAGCA	1200
	GATGCCACCG CCAGAAGAGC CCTTATGACC CCTGGGCCGC GCTGTGCCTT AGTGTCTACT	1260
50	TOCAGGACCC TTCCTCCTTC CCTAGGGCTG CAGGGCCTGT CCACAGCTCC TGTGGGGGTG	1320
50	GAGGAGACTC CTCTGGAGAA GGGTGAGAAG GTGGAGGCTA TGCTTTGGGG GGACAGGCCA	1380
	GAATGAAGTC CTGGGTCAGG AGCCCAGCTG GCTGGGCCCA GCTGCCTATG TAAGGCCTTC	1440
5 5	TAGTTTGTTC TGAGACCCCC ACCCCACGAA CCAAATCCAA ATAAAGTGAC ATTCCCAAAA	1500
	AAAAAAAAA AAAAAAAAA ANCCCGNGGG GGG	1533

	(2) INFORMATION FOR SEQ ID NO: 91:	
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 575 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 91:	
	ATCCTCTGGA ATCTAGGTGG AAGCCACCAA GCCTTCTTCA CACTTGCGTT CTGAGCATCT	6
15	GCAGACTTAA CCCCATGTGG CAATCACCAA GGCTTATGGC TTGTGTCCTC CAGAACTGTG	12
13	GCCAGAGCTG TACCTGGGCC CCTTTGAGCT GAGGCTGAAG CCAGAGTCTG AAGCTCAGCA	18
	GGGCAGTARG GCCCTGGGCC TGGCCCCTGA AACCATTCTT TTCTCCTAAG CCTCTGGGCC	24
20	TTTGATGGGA RGGCTGTCC TCAAGATTTT TGAAATGCCT TTGGAGGGTT TTTGCCTTGT	30
	CTTGGATATT GGCTTCCTTT TAGTTATGCT CATCTCTCTA GCAAGTGAAT GTTTCACAAC	36
25	CTGCTTGGAT TCTTTCTCTA CCACAGARCC AGGCTGCAAA TTTTACAAAC TTTTACACTC	42
23	TGTTTCCCTT TTAAATATAA ATTTCAATGT TAAGTCACTT CTTTGCTCCC ATATCTGATT	48
	TAGGITGCTG GAAGTAGCCA AGTCACCTCT TGAATGCTTT GCTGCTTAGA AATTTCCTCT	54
30	ACTAGGTAGC CTGGGTCATC ACACTTAAGT TCAAA	57
35	(2) INFORMATION FOR SEQ ID NO: 92: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 639 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 92:	
45	TCCTTCATC TTAAGCACCA CCCGACAGGG CAGGTACTAT TACCATCTCC GTTTGACAGA	6
	TNAGGAACCT GGCACAGGAA GCATTTAAGT GGATTCCCCA GGATCGCCCC ACTGTCAGGA	12
50	GCAGANTCAG AATGGGCCTC AGCATCAGGC TCCCAATCCT GGCTTCTAAC TGCTGCGCTC	18
	TGCCCTTCYC TCWCCCCACC TCCCCACTCC AGTGCCTTTG GTCATGCCAC TGCAGCTTTC .	24
E =	AGGCCAATAC TOGATTAGCC TCTTAGTGTT CTTGTCCCTG CAGCCATTTC CCCAGGCAGC	30
55	AATTCCATGT GCCCTCACTG ATGTAGGTGG CTCTTGTGTC ATTTGTCACA TCCTATTGAA	36
	TTGTTTATGC ATCTTGTTCA CACTCACAGC ACCCTCCCTC TCACACGTCC TCCTTATAAA	42
60	AATGTCCCTC AGTGTCTGCT ATGAGCCAGG TGCAGACTTA AGTGACAGGG CTGCTACGGG	48

	AAATAAAAAA TTAACAAGGA GCACCTGCCT CTTAATGCAC AGTAACAAAC TATGTTAAGT	540
	GTCAGGAAGG AAAGGTTAAG GATGCCAGGA AGGCTTTTAA TAAATAACCT GACTTAGATG	600
5	GGCAGGTGGT GCTGARGATT AAGAACGTGT TCTTCTCGA	639
10	(2) INFORMATION FOR SEQ ID NO: 93:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 744 base pairs (B) TYPE: nucleic acid	
15	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:	
20	GAATTCGGCA CGAGAGTGGC TGGAGTCTGG CTGCAGAGGG AAGACATCAG CAGGGAGGGA	60
	GCCAGGGCCT GTCACATCTT TCCTCTGGCC ATTGTCCTGG TCTTTGTAAG CCCAGAATCT	120
25	CCCCTTCCCT GAAGGGAGGC CAGCACCCCA GGAGGGCAGC AGGTGTGCTG TGAGGGTTGG	180
	AGTAGTGTGA GAGGTCAGGG TACACTAGAA TGGCCATGGA CACCATGTGG GGGTGCTCTG	240
	GOCTGGGCCA CAGAACAGTG TCCTTCCTGC TGCTCCTCCC CTGCAGCTTC CCCCGACCTT	300
30	GINGITTATT TOGTTTGATA CCAATCAGCA GACCCTGCAA GGTGGAAGCT CCCAGGCTCT	360
	CAGTCCCACS ACTCTCATGT GCCAGTCACC CNTACTGTAA CTGCCCAATG AGTACTTCTT	420
25	GCCCACTGCC AAGATAGAGC CAGTTTACCA AGACAGGGGA ATTGCAGTAG AGAAAGAGTT	480
35	GAATATACAT AGAGCCAGCT AAATGGGAGA GTGGAGTTTT CTTATTACTT AAATCAGCCT	540
	CCCYTAAAAT TCAGAGGTGA GAATTTTTCA AGGACAGTTT GGTGGSCAGG CCTAGGGAAT	600
40	GGATGCTGCT GATTGGCTAG GGATGCAATC ATAGGGGTGT AGAAAAGTWC CTTGTGCACT	660
	GAGTCCACTT TTGGTGAGAG CTACCAAGGA GCTGCTGGTC TGCTGGTCCC GGTAGAGCCA	720
40	TCTGGTGTCA GGAATGCAAA AGTG	744
45		
50	(2) INFORMATION FOR SEQ ID NO: 94:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 526 base pairs (B) TYPE: nucleic acid	
55	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 94:	
	GCAGGGGAAT TCGGCCACGG AGGGGTTTCA ACAGGGCCCG TGGGGTGAGG TGCARACACA	60
60	The state of the s	, ,

	AAGCCCATAA GTGCTGGCCT GTTGGGACAA ATGAGAGAAA TCCCATAGGG TGGTGATGAC	120
	AGCGCAYTCA GCCATCYTAY TCCTGGGGAA AATGAAACTT GTGCTCCTAT CAAATGCTCA	180
5	GTTGTAAAAC ŢGGAAAAAAA TTTTAGAAGA CATCTTGTCC AGCATCTGTG TTTATGTCTA	240
	TAAAATGTAG AAAACTAAAG CACAGAGATG TTAAATGTTT TGTCCAAGGT CCAACAGCTG	300
10	GTTAGCARGC TTGGTCTGGT GACCTTTCTA CTGAACCACA GTGCCGCTGG GGGAAGTCCT	360
10	CAGCACAGAT GOCTGCTG ATAGCTGGGG TATGGGCAGT ATTAGTAGTT AACCAGTCAA	420
	CCCAAGTTCC CATAGTCTAG GTTCTGCTTC AGCTGGAGGT TAGGGAAAAA CACAAGAAAA	480
15	TCCCTTACCA CTCTACCAGT GCTGGGGGAT GTACTAAGAG ATCCCC	526
20	(2) INFORMATION FOR SEQ ID NO: 95:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 426 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 95:	
30	GGCACAGGGC AGGAGAGACT TGGTCCATGG GGAGAAGCCT GCAGTATAGA TGGGACCTCC	60
	AGGAGCCCAA GTAGCATAGA CCCTGCTGAT CCGGGGCCAT TGAGCCAGAG GATTTGGGCT	120
35	GAATGTCCCC AGAGACAAAA GOGAAAGGTA GATCCTTTCC CTTAAAGATG AAAGCCATCG	180
	CCCGGGCTTG CTTATTGCTC TCTCTCCTGG TCCTTCCACA TGTTGTTTCT GAACATTTGT	240
	TCTGGCATCA CAATCCCCGT CATCCTGTCA TCTGGCCCTT CCCACCTTTC CACCTTATCT	300
40	CTTGCAGTGT CTCCGCGTCG ACCTGGCACC TGGGTGAARG CTTGCTCTTG CTGGTGCCCA	360
	TAGCCCCCAG TGTATGGTCT TGAMCTCCCC AGCCATATGG ARACCCACCT CAGGAGGGCC	420
45	CCTCGA	426
-1.5		
	(2) TITOMORTON POR CTO TO 170 06	
50	(2) INFORMATION FOR SEQ ID NO: 96:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 844 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:	
60	GGCACAGCGG CACGAGATAG GAAGCTTGGC AGGGGCAGCT CCCCCAGTGC GCATTGCCCT	60

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•	GTAACTCGAG CGCCTGGGAG TGGGGAGAGG CTTGGAAATG GAGCAGGGTG GTGGACCTCG	120
	TOTTOTOCTG CTCATCCCAG GCCTCCTCCA TAACACCTAC CTAGCACGGC CTGGGGACTT	180
5	CCCAGCCCAA GGAACAACTG AGAATACTGA GTGCCAGGGT AGCCCTAGCC CCATTTCACA	240
	CCTGGGCAAA GTGAGGTCAC TGGATTCAAA CACTCAGATT TAAACCTCCT CTGTGTCTGC	300
10	AGCACCTGTA TATAACTGCC AGCCTCTGCT GCCCCTCTCC AAAAAGTCTC TGCCCTTGTC	360
10	TTTGGCACCT GTCTCTGTCC TCCCCATTCT CTGCTCCTCC TTTCTCCAAC TCAGANTCAC	420
	CCTGTTAGTT CAGCAAATGT TCATCGAGCT CCATAATGTA GCAGGACAGG NCTGTCTAAC	480
15	AGATTCTGGN CTTGCAAGGG TGAGACAAGT ACTCTCCATC TTTCTCTCAT CTTCACAGAT	540
	GGTCTGCTCA ACAACTTTGC ACTGAATTGT AAATAATTGA TACTGCATAA AACATTGATG	600
20	TTCTTTAAGG GTAGTCCAGC AAGGTGGCAA GTCTTATAAT GATAACTGCT CAAGGATCTC	660
20	TCAGTGAAGC ATTTGGGGST GCTAGCTCTG CCTATGGGTG AGGTCAGCTA TCTCACGCCA	720
	TCTACTTCCA CNTGCCCCCC CATGCCAGGC TCACCCTGAG CTGAGATGCC TGAGCAGGTG	780
25	GCAGAAAGGA GCCACCTGGT TTATGCTTCG GGACCACAAA CTCCTCTATC CAGANGACAG	840
	TTTT	844
30		
	(2) INFORMATION FOR SEQ ID NO: 97:	
	(i) SEQUENCE CHARACTERISTICS:	
35	(A) LENGTH: 1985 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(b) received. Italical	
	(xi) SECHENCE DESCRIPTION: SEC TO NO: 97:	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 97: ACCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC	60
٠.	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC	60 120
45	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG	120
45	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT	120 180
45	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG	120 180 240
	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC	120 180 240 300
50	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC CCCCATCCTG GTGCCTCACA CAGCGCAGCG GNAGAGCAGA GGTATCCATT TGAGGCCCTC	120 180 240 300 360
	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC CCCCATCCTG GTGCCTCACA CAGCGCAGCG GNAGAGCAGA GGTATCCATT TGAGGCCCTC TTCCGCAGCC AGCACTACGS CCTCCTAGAC AATTCCTGCC GCGAATACCT TTTCATCTGT	120 180 240 300 360 420
50	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC CCCCATCCTG GTGCCTCACA CAGCGCAGCG GNAGAGCAGA GGTATCCATT TGAGGCCCTC TTCCGCAGCC AGCACTACGS CCTCCTAGAC AATTCCTGCC GCGAATACCT TTTCATCTGT GAATTTTTTG TTGTGTCTGG CCCAGYTGCA CACGACCTGT TCCATGCTGT CATGGGCCGT	120 180 240 300 360 420 480
50	AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG CTCTTACCTG GGGCGGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC CCCCATCCTG GTGCCTCACA CAGCGCAGCG GNAGAGCAGA GGTATCCATT TGAGGCCCTC TTCCGCAGCC AGCACTACGS CCTCCTAGAC AATTCCTGCC GCGAATACCT TTTCATCTGT	120 180 240 300 360 420

	GTTCCTGCCC	TOGACAGGTA	CTGGGGAACA	GGTGCTTGCC	TTGCTATGGC	CACGGTTTGA	660
5	ACTGATCCTG	GAGATGAATG	TTCAGAGCGT	CCGAAGCACT	GACCCCCAGC	GCCTAGGGGG	720
J	GTTGGATACT	CGGCCCCACT	ATATCACACG	CCGCTATGCA	GAGTTCTCCT	CCCCTCTTGT	780
	CAGTATCAAC	CAGACAATTC	CTAATGAACG	GACCATGCAA	TTGCTGGGAC	AGCTGCAGGT	. 840
10	GGAGGTGGAG	AATTTTGTCC	TCCGAGTGGC	AGCTGAGTTC	TCCTCAAGGA	AGGAGCAGCT	900
	TGTGTTTCTG	ATCAACAACT	ATGACATGAT	CCTGGGTGTG	CTGATGGAGC	GGGCTGCAGA	960
15	TGACAGCAAA	GAGGTTGAGA	GCTTCCAGCA	GCTGCTCAAT	GCTCGGACAC	AGGAATTCAT	1020
13	TGAAGAGTTG	CTGTCTCCCC	CTTTTGGGGG	TTTAGTGGCA	TTTGTGAAGG	AGGCTGAGGC	1080
	TTTGATTGAG	CGTGGACAGG	CTGAGCGACT	TCGAGGGGAA	GAAGCCCGGG	TAACTCAGCT	1140
20	GATCCGTGGC	TTTGGTAGTT	CCTGGAAATC	ATCAGTGGAA	TCTCTGAGTC	AGGATGTAAT	1200
	GCGGAGTTTC	ACCAACTTCA	GAAATGGCAC	CAGTATCATT	CAGGGAGCGC	TGACCCAGCT	1260
25	GATCCAGCTC	TATCATCGCT	TCCACCGGGT	GCTGTCCCAG	CCGCAGCTCC	GAGCCCTCCC	1320
	TGCCCGGGCT	GAGCTCATCA	ACATTCACCA	CCTTATGGTG	GAGCTCAAGA	AGCATAAGCC	1380
	CAACTTCTGA	TGTGCCAGAA	ACCGCCCTGA	GATCTGCCGG	TCATCTCCAT	GGACTTCTGC	1440
30	ACCCCATTCC	ATACCCTTCT	TCACCTGGGG	TACCCCTTCC	AGTTTTCCCC	TTGCTTCCCA	1500
	GGCCCTTGAC	ATGGCTTACC	TGCCTTCACT	CCCAGCACCT	TGCCCAACAG	GATAAGCTGG	1560
35	ATCCCCTTGG	CCTTCTGAAT	ATCCCAGTGT	CTTCAGGTTT	CCCAAGACCA	CTTCCCTGTG	1620
	GCCTTCCAAA	ATGGCCTTTA	TCATTTCTCC	AGTCTGTCAC	CCTCCTTTCC	TGCTCCCATA	1680
	CACCCAAGGC	TIGITICITC	CCCTGTAAAA	ACCACTGCCT	CAATCTCTGG	TTCACTCAAC	1740
40	TAGTCACCAT	GTCCTGAGGC	ATGAAGCCTC	CTCAGCTCTT	GGAATTGCTG	GCAAGGGGTG	1800
	ACTGCCTCTG	AGTCATTGTG	TTTTTCAAAG	TGATTTCTTT	TCTGTAGCTT	TTTGACCTAA	1,860
45	GATCTCAGCA	ATTTGAACAC	TAACCTCTCC	CCTCCTGGCT	CAAGAATTAC	TCCGAAGTCA	1920
	GTCTGCAGAA	AATAAATATT	TAGTATGACA	TGAAAAAAA	алалалала	АААААААА	1980
	ААААА						1989

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(2) INFORMATION FOR SEQ ID NO: 98:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1416 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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1416

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 98:

	ATATGAAGGG	AAAGAATTTG	ATTATGTTT	CTCAATTGAT	GTCAATGAAG	GTGGACCATC	60
5	ATATAAATTG	CCATATAATA	CCAGTGATGA	CCCTTGGTTA	ACTGCATACA	ACTTCTTACA	120
	GAAGAATGAT	TTGAATCCTA	TGTTTCTGGA	TCAAGTAGCT	ATTTATTA	TTGATAACAC	180
10	AAAAGGTCAA	ATGTTGGGAC	TTGGGAATCC	CAGCTTTTCA	GATCCATTTA	CAGGTGGTGG	. 240
10	TOGGTATGTT	CCGGGCTCTT	CGGGATCTTC	TAACACACŤA	CCCACAGCAG	ATCCTTTTAC	300
	AGGTGCTGGT	CGTTATGTAC	CAGGTTCTGC	AAGTATGGGA	ACTACCATGG	CCGGAGTTGA	360
15	TCCATTTACA	GGGAATAGTG	CCTACCGATC	AGCTGCATCT	AAAACAATGA	ATATTTATTT	420
	СССТАААААА	GAGGCTGTCA	CATTTGACCA	AGCAAACCCT	ACACAAATAT	TAGGTAAACT	480
20	GAAGGAACTT	AATGGAACTG	CACCTGAAGA	GAAGAAGTTA	ACTGAGGATG	ACTTGATACT	540
20	TCTTGAGAAG	ATACTGTCTC	TAATATGTAA	TAGTTCTTCA	GAAAAACCCA	CAGTCCAGCA	600
	ACTTCAGATT	TTGTGGAAAG	CTATTAACTG	TCCTGAAGAT	ATTGTCTTTC	CTGCACTTGA	660
25	CATTCTTCGG	TTGTCAATTA	AACACCCCAG	TGTGAATGAG	AACTTCTGCA	ATGAAAAGGA	720
	AGGGGCTCAG	TTCAGCAGTC	ATCTTATCAA	TCTTCTGAAC	CCTAAAGGAA	AGCCAGCAAA	780
30	CCAGCTGCTT	GCTCTCAGGA	CTTTTTGCAA	TIGITTIGIT	GCCAGGCAG	GACAAAAACT	840
50	CATGATGTCC	CAGAGGGAAT	CACTGATGTC	CCATGCAATA	GAACTGAAAT	CAGGGAGCAA	900
	TAAGAACATT	CACATTGCTC	TGGCTACATT	GGCCCTGAAC	TATTCTGTTT	GTTTTCATAA	960
35	AGACCATAAC	ATTGAAGGGA	AAGCCCAATG	TTTGTCACTA	ATTAGCACAA	TCTTGGAAGT	1020
	AGTACAAGAC	CTAGAAGCCA	CTTTTAGACT	TCTTGTGGCT	CTTGGAACAC	TTATCAGTGA	1080
40	TGATTCAAAT	GCTGTACAAT	TAGCCAAGTC	TTTAGGTGTT	GATTCTCAAA	TAAAAAAGTA	1140
+∪	TTCCTCAGTA	TCAGAACCAG	CTAAAGTAAG	TGAATGCTGT	AGATTTATCC	TAAATTIGCT	1200

(2) INFORMATION FOR SEQ ID NO: 99:

45

50

55

60

(i) SEQUENCE CHARACTERISTICS:

AAAAAAAAA AAAAGGAAAC TCGAGGGGG GCCCGG

(A) LENGTH: 1935 base pairs

GTAGCAGTGG GGAAGAGGGA CGGATATTTT TAATTGATTA GTGTTTTTTT CCTCACATTT

GACATGACTG ATAACAGATA ATTAAAAAAA GAGAATACGG TGGATTAAGT AAAATTTTAC

ATCTTGTAAA GTGGTGGGGA GGGGAAACAG AAATAAAATT TTTGCACTGC TGAAAAAAAA

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 99:

5	NICTACCCTA ATCAAGATGG GGACATACTT CGCGACCAGG TTCTTCATGA ACATATCCAG	60
3	AGATTGTCTA AAGTAGTGAC TGCAAATCAC AGAGCTCTTC AGATACCAGA GGTTTATCTT	120
	CGAGAAGCAC CATGGCCATC TGCACAATCA GAAATCAGGA CAATAAGTGC TTATAAAACC	180
10	CCCCGGGACA AAGTGCAGTG CATCCTGAGA ATGTGCTCTA CGATTATGAA CCTCCTGAGC	240
	CTGGCCAATG AGGACTCTGT CCCTGGAGCG GATGACTTTG TTCCTGTGTT GGTGTTTGTG	300
15	TTGATAAAGG CAAATCCACC CTGTTTGCTG TCTACTGTGC AGTATATCAG TAGCTTTTAT	360
15	GCTAGCTGTC TGTCTGGAGA GGAGTCCTAT TGGTGGATGC AGTTCACAGC AGCAGTAGAA	420
	TTCATTAAAA CCATCGATGA CCGAAAGTGA CCAAGACCAA GGCCCACCAA GGCAGCAGAC	480
20	TGTTAATCAG ACAAACAGAT CTCTGAGAAG GTGCATCAGC TGCTTTGAAG GCTGAAGATT	540
	GTTTTGTATG ATACTGCACA GCATCAGGCA TTTTAAAGCA GATCTTTACT AAACAGGTTA	600
25	ATGAGCTAAC AAGCAGGITC TCTCGTCTTT GGGCTCTTTC CTTTCTGAGT TGCATATTCT	660
23	ATTITICTTGT CCCCAAGTAG AGACTAGTAC TACAAAAAGG GACCACATTT TTCAAGTATT	720
	TCTAAGTATA AAAAACAAAA CAAAAATCTC TTAGGAAATG TCTAGACCTC CATTCTTGGA	780
30	TTCCCTTTCT TTCCTTTTAT TTTAAAAAAG AACAGTACCC CTCTTTTAAG ATGCTGTCTT	840
	ACATTAATGA GCATCTAATG GAAAGAAGGT ATGAGTTGCA CTGAGGATTA GAATAGTGGT	900
35	GCGTTAGTGG CATTATCTAT AAATACACTC ACCTAAATTG AAAGCTAAGA AGGAAATGTA	960
))	AATATAATAT ATATTTATAT TIGATGTAAT ATGGACATCT GCAGATTCTA ATAAACAAGG	1020
	ACTATTCCTG ATAGTAGGCT GTGACATACT GTCTTGTGAA ATGGTTTCCT TGACAAAATT	1080
40	TAAGCTGAGC TTAAAAGCAA AAAAACAAAA AGTACACAGA AATATTTATT AAAATGTAAT	1140
	ACAGTTTATT GAACTTTCTA GGTATGGAGT TTGATGGACA GGGCTGCCTY TAATGAGTGT	1200
45	GAAGGTCACT AAGTCACTTA GACATCTCAC CGTGGAAGTT TGTGAGCCTG CATTAGGAGA	1260
73	TAGACTGATT ACCATACATG ACATAAAAAG GAACAGTGGA TAGCTCATAC TTTATGGTGG	1320
	TTCTTCTCCT CCGAAATAAT ATACTGCAGA AATCCCAGAC AGAGCTCCTF ACAAACCTTT	1380
50	AATTGTAATA TATTTTGAT GATTATTCAC ATTGAATGCA CAGACCAAGA ATTCAGTGAA	1440
	TGTCATTTTT TAAAAAACTA ATTTGTATTG TCTGCTCTAG TGATACAAGT TTTACTAGTG	1500
55	ATAAACTATT TTAATCAACC ATACTATTCT TATGGAAAAA AATATCTATT TTGGCAGGTT	1560
<i>J J</i>	TCTGTGCCTT TATTTCCCTC TTCTGAAAAA AAGTCTGTGT TTTCATAGTT TGGTTTGCAT	1620
	TGTATATCAA TAATTAATCA GGAATGGGTT TTGGTGCCTG AAAAATTGGC CATGGAGGCA	1680
60	CACCAAAGCT TCAAGCACAA GTCTTGTACA TGGGCCATCA CTGTCTGGTT TCACTTCGTG	1740

	TGTTTCCTAA ACACATTTAG CTGCTTTTTT AACAAACTCA GCCCCATACT TGAGTCCCTT	1800
5	GTTGTTGGGA GCATTTCCAG GCATCTTTTA AGGGAACTGT GACAAACAGC CTCGGGCAGA	1860
J	TGAACACGGA GCCTCTCTGT TGTCTGTCTC TGAGATCTTT GTGTCTGGGA ATGCCTAAAG	1920
	MITTEMITT TITT	1935
10		
	(2) INFORMATION FOR SEQ ID NO: 100:	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 599 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
••	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 100:	
	GAATTCGGCA CGAGCGTCCA CGCAGCCGCC GGCCGGCCAG CACCCAGGGC CCTGCATGCC	60
25	AGGTCGTTGG AGGTGGCAGC GAGACATGCA CCCGGCCCGG	120
	CCTCATCCTG ATGGGCACTG AACTCACTCA AGACTCCGCT GCCCCCGACT CCCTGCTGAG	180
20	AAGTTCAAAG GGCAGCACGA GGGGGTCTTT GGCTGCTATT GTCATCTGGA GGGGGAAGAG	240
30	TGAGAGCCGG ATAGCCAAGA CCCCAGGCAT TTTCAGAGGT GGCGGGACCT TAGTCCTACC	300
	CCCAACACAC ACCCCTGAGT GOCTCATCCT CCCTTTGGGC ATAACGCTGC CCTTGGGGGC	360
35	TCCAGAAACA GGCGGTGGGG ATTGTGCCGC TGAGACCTGG AAGGGCAGCC AGCGTGCCGG	420
	CCAGCTGTGT GCATTGCTGG CTTAATATGC AGGCTTGGG GGGCTGTGGC CACATGCCCG	480
40	GCAGGAGGTG AGTGAGGAGC CCTGTGGCGT GCTGGTGTGG GGATCGTGGG CATTTCAAAC	540
	GGGCTTGTCG TACCCTGAAC AATGTATCAA TAGAGAAAAA AAAAAAAAAA	599
45	(2) INFORMATION FOR SEQ ID NO: 101:	
	•	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 784 base pairs	•
50	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 101:	
<i></i>	GAATTCGGCA CAGAAAAAA AGAGAGACTG GGTCTTACTG TGTTGCCCAG ACTTGTCTTG	60
	AACTCCTGCC TCAGCCTCTC AAGTACTTGG GATTATAGGC CAAGAAGCCA CCATGCCTAG	120
60	CHALLIANCE ACTUALIST SESCULTABLE ACCOUNTS CALLETAIN ALLIANTED	180

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	CACTITISCAC ATCCACTIGIT CACCAAATCK RGITICATICIT GCATCCTAAG TAAGTCCTTT .	240
5	GATTCCTCCA GITGITCATT AGTAATGTCT CAARTGTAAT TITTTCTAGT AGTITTCAGC	300
J	CTGTCTTTCC KGCCTTCAGT CTTAACTTCT CCAGTACATA KGCCACATTG TTGTCAGCAK	360
	GATCAWATTT TATTTAAAAA TACTTTACAW AKGTTTATKG CCAAATATTA GRAAATACAG	420
10	ATTCATGGAA AGAAAATCA CTGTCCCAAG GAGGTCACTG GCATGGTGAG GTTAAGGGGT	480
	GATTITAATT TITAAAAATG TATATTITIT CCTGTGTAGA GTAGTAACAC CCTTGAAAAC	540
15	ACAMICCCTT GTAAAGTCTC TAATTCTGTA CTCCGCATCT AGSTGRTCTC TTCTTTCTCA	600
15	GATATTITAC AATTICATTI ATCACCACCT TICTCTAGCC TITACCCGTC TCTTCAATAT	660
	TWACATATGC AGAAGTITCT CCTAACAAAC ACCTGCCTCT GCCTCAGTTC TGCTACCACC	720
20	CTGTTGCTTT CTTTCCCTTC ACAATCAAAT TTAAGAGTGT CAAAAAAAAA AAAAAAAAAA	780
	TOGA	784
25		
23	(2) INFORMATION FOR SEQ ID NO: 102:	
30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1035 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:	
	AGAGGCCTGG CTGCGTTGCC CTATCTCCGT CTCCGCCACC CACTTAGCGT TTTAGGCATC	60
40	AATTACCAGC AGTTTCTCCG CCACTATCTG GAAAATTACC CGATTGCTCC CGGCAGAATA	120
	CAAGAGCTTG AAGAACGCCG CAGTTGCGTG GAAGCCTGCA GAGCAAGGGA AGCAGCGTTT	180
	GATGCCGAAT ATCAGCGAAA TCCTCACAGG GTGGACCTCG ATATTTTAAC CTTTACGATA	240
45	GCTCTGACTG CCTCTGAAGT TATCAACCCT CTGATAGAAG AACTTGGTTG CGATAAGTTT	300
	ATCAATAGAG AATAGTTAGG TGGTGACACT ACTTCAAGAG AACCTCTGCA TTCCAGTCAT	360
50	ACCAATCCTG CAACTTGATT TTCAGAAGTC AAGAGTATAT CGCGATAAGA CAGTGCACAG	420
	GTGGAGGGGA AAAAAAGGGG GAGGGGGAAG CTTATCTTGA AAAAGCATCA CAGAAGTAGA	480
	AAAAAATGTC GAAAGCATTA TAACTGTAAC GFTCTTTGAG TTTGTGATTG ATCCACATTT	540
55	TICCCCCTGC ATTATGGAAA ATGTCTCTCA GCATTGCTTT ATTACAAAGT AAAGGATGGT	600
	TTTATAAAAT TGAGACTGAT GAAACATCAA TACTAGAGCC CATGAGGATG AAAGAAATTA	660
	TCAAATAGTG CTGAACAGAA TAAGATGTTA ACGCTGAGTT ATTAGGACTG GAAGGCTATG	720
60		

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	AAAAGAACTT GAAATTGTCG GAATATGTGC TCTCTTCATG TCATATTCAA TAGAAGTTTC	700
		780
_	TAGTTTAAGA TIGATTTTGT GITTTCTTAG GCATTTCAAG TGACAAGCAA AGTAAATGTA	840
5	TATATTATGT GATAAATCAT GTTTTCAAGA ACGTCAAATT TCTGGACTTT TTTCTTTCAA	900
	TTTTTAATIT TTAAAGTTTT TTTGGTATTA AAAAATCYAT TCACAAGCCA AAAAATWTWT	960
10	WAAATWIWCM GCGAAAAGCC AAAAAAAAAA AAAAMMAGGG GGGGCCGGGC CCCATCCCCC	1020
•	CAAGGGGGTC CNGNT	1035
15	(0)	
	(2) INFORMATION FOR SEQ ID NO: 103:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2218 base pairs	
20	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103:	
	AGGTATTAGG CCCTTTTGTG GGAGCCCCAT GTTTTGTTTT	60
	SGGAGGGGA GGCCTGAATT GTTTTGCAGA GGAAGATGGC ATCTGTGCTT TAAATTTCTC	120
30	ATTACTOGGT TAGAAAACAA AGAGGGAKTG CCCTGCACAT TITCTTTTGT GCTTTTAAAT	180
	GTTTCTTAAG TTGGAACAGG TTTCCTCGGG CCTGTTTTGA CTGATTGCTG GAGTGCATTT	240
35	GATAGTTAAA AATTACTAAT TGGTTTTATT TCCCTTCACA CTCTGCCTCC CCACTTCTCC	300
,,,	CCCCGTTACT GAAAAATAAC CATTTTAGTG TCAGGCTAGA AATTGAATTG	360
	TGTATCCTTT AAATTAAAAA CCACAAGTGT TTATTGTAGT GGTTAAACTG TAGCATCTCA	420
40	GCATCTGGGT GGAAGCTGCC TATATTTCTT CCCAGTTTAA CTGGGGACCA TCTGTGAAAT	480
,	TAATTTTCCA TCCAGACAGC TGCTGTGAGC AAATGAACAT AAATGCTCGC TGGAAATTTA	540
45	CTAACCAGTT TTTATATTGA CCTGCAGTGT AAAAAGCACA TTTAATTATA AACAATATAT	600
	TCAAAATGGG CAAATTTTAT TTTCAAATGC AGTGTAGAGC TAGATTAAAA GCAACTCTTT	660
	GCCACCTACT CTGCCCTTTT GGCAAAGTTA CCTTGAACAA AGAATCTTAA GGGTTTATTA	720
50	AGAACTCTTT ATTTTCTTCA TACCCTGTTC TCTGCAGTGC TTTCTAACAG CTTCTGGGTG	780
	CAGATTITCT TCGGCATCCT TITGCACTCA GCTTATTACA GGTAGGTAGT GCTTAAGAAA	840
55	AGTCATGGAG GACTAAAGCC TAAGTCCTTT TCACTTTTCC TCCATCTGAA GGTAGGTGAG	900
JJ	TTCATCCTCT TCATAGTAAT GCTGTTTTAC CAAGACTTTA TAGCAGATGG ACCCAGAAAG	960
	AATTTTCTGC TATTGTGTTC ACTACAACAG GATAGGGACA TCAGACAGCC CCAGAAACCC	1020
60	CTTCCAGATC TGATATGGGA CTATTAATTT TTATGCTGTT AATTGGTATT CATTCACAAT	1080

	GCAGTTGAAG	GGGGAAGGCT	CCACTGCATT	CTTTGGCTAA	GGCCTGAATG	CTTGCTCATC	1140
5	TGTAAGATCT	ATACTCGAGG	TTITGTTTC	CTTTTAAAAT	TCTTTAGGGA	GAGAGGGATG	1200
J	GTTTCTGAGG	GGTTCTGAAA	GTATGATTCA	ATGTGCAACA	TACAGGTAGG	TCTTCAGCAT	1260
	AAGCTGAAAT	ATATGCATGT	AAAAACTTTG	ACATCTTTTT	TTTTAATTTT	CCACTTTCTT	. 1320
10	CTTAACTTTA	CTTCTCTTTT	TGTCCCCCCC	CCATCTTACA	GAAGTTGAGG	CCAAGGGAGA	1380
	ATGGTAGGCA	CAGAAGAAAC	ATGGCAAACT	GCTCTGTGCT	TTCAAACCAA	AGTGTTCCCC	1440
15	CCAACCCCAA	ATTIGICTAA	GCACTGGCCA	GICIGITGIG	GGCATTGTTT	TCTACAACCA	1500
15	AATTCTGGGT	TTTTTTCTTC	TTTCTTTAAA	CATAGAGGTA	CCACCACAAG	GGATGCCCTA	1560
	CTCTCTCGCA	GCTCTTGAAA	GCATCTGTTT	GAGGGAAAGG	TCTCTGGGCA	AGCAAGTGGT	1620
20	TATTTGGATT	GCTTGCTTCC	CTTTTTCCAC	CTGGGACATT	GYAATCATAA	AATAACAGTA	1680
	AATTCCAAAC	CTCAAAAACT	ATTATGGCCT	GAGCACAGCT	GAAATCTAGC	AGAGTTTAAC	1740
25	TCTTCTGCCT	CCATGTCTGT	CACTTATAAT	TCAGGTTCTG	CTGTTGGCTT	CAGAACATGA	1800
23	GCAGAAGAAT	CGTTTTATGC	TAGTTATTGC	ATTCATGGTT	GAAACTCAAC	TTAGGGAAAG	1860
	GGTTCCAATG	TATTAAGCAA	TOGGCTGCTT	CTCCCCAATC	CTCCCTAACA	ATTCGTTGTG	1920
30	TGGACTTCTC	ATCTAAAAGG	TTAGTGGCTT	TTGCTTGGGA	TCAGTGCTCT	CTATTGATGT	1980
	TCTTGCTGGT	CTCCAGACAC	ATTCCTGTTG	CATTAAGACT	TGAAAGACTT	GTAGATGTGT	2040
35	GATGTTCAGG	CACAGGATGC	TGAAAGCTAT	GTTACTATTC	TTAGTTTGTA	AATTGTCCTT	2100
<i>JJ</i>	TTGATACCAT	CATCTTGTTT	TCTTTTTGTA	GGTATAAATA	AAAACACTGT	TGACAATAAA	2160
	алалалала	АААААААА	АААААААА	ааааааааа	АААААААА	ААААААА	2218
40							
	(0)						
	(2) INFORM	ATION FOR S	EQ ID NO: 10	04:			
45	(i)	(B) TYP	HARACTERIST GTH: 1351 b E: nucleic ANDEDNESS:	ase pairs acid			
50			OLOGY: line				
50	(xi) SEQUENCE	DESCRIPTION	: SEQ ID NO	: 104:		
	CTTCACAGAC	TGACAGAATG	GTTTGTTT	CITTICITIT	CTTTTCTTTT	GTTTTTGAGA	60
55	TGGACTCTAG	CTCTGTCACC	CAGGCTGGAG	TGCAGTGGTG	CGATCTCGGC	TCACTGCAAG	120
	CTCCGCCTCC	CGGGTTCTCA	CCATTCTCCT	GCCTCAGCCT	CCCGAGTAGC	TGGGACTACA	180
	GGCGCCCACC	ACCACGCCCG	GCTAATTTTT	TGTATTTTT	AGTAGAGACG	GGGTTTCACC	240
60							

	ATGTTAGCCA GGATGGTCTC GATCTCCTGA CCTCGTGATC CGCCCGCYTC GGCCTCCCAA	300
	AGTGCTGGGA TTACAGGCGT GAGCCACCGT GCCTGCCCCA GAATGGTTTT TAAAGCCACA	360
5	GTTGAGARGC CACCCATTGC CCGGCGCCTG GACAGTGATC ATCTTGTTCA TCTTGTTCAG	420
	TCCTTTCTTG TGTGATTGGA ATTATTCATC CCCTTTGAAA GATGAGAAGG TTGAGATGCA	480
10	AAGAGTCTAC CTTTCCAAGT TCTCACTGCT GGAAAGARCT AGAAGCACAG TTCAAAGTTC	540
	TOGNITICTOG ACTICTOCAGT CCAGGTYTCC CTTYTCCCAC TTGCCTACCC TCAATGCCAC	600
	ACTGTTTTG AAGTGGCCCA TAACTTGAAG GRAAAGTTTA AAGACAGTTC AATTTAATCA	660
15	TCAGRATGCA TTCTTTTTT TTTCGGARAC GGAKTTTCAC TCTTGCTGCC CASGCTGGAG	720
	TGCAATGGTG CAATGATCTC GGCTCACTGC AACCTATGCC TCCTGGGTTC AAGNGATTAT	780
20	CCAGCCTCAG CCTCCCGAGT AGCTGGGATT ATGGGCGCCC ACCACCATGC CCAGCTAATT	840
20	TITGTATTIT TITTITIAGT AGAGATGGGG TITCGCCAGG TIGGCCAGGC TGKTCTIGTG	900
	AAYTCCTGGC YTCAGGTGAT YTGCCCACYT CATCYTCCAA AAGTGCTGGG ATTACAGGCA	960
25	TGAGCCACTG CGCCTGGCYT CAGAATGCAT TCTTACACAT CTATCCTAGA CATTTATAAG	1020
	CACTCTAATG GATAACAATC CAAGAATAAA TGATTGTAAA AGATGATGCC GAAGAGTTGA	1080
30	TGTCAATCTT TTTTTCCTAA GAAAAAAAGT CCGCGAGTAT TAAATATTTA GATCAATGTT	1140
30	TATAAAATGA TTACTITGTA TATCTCATTA TTCCTATTTT GGAATAAAAA CTGACCTTCT	1200
	TTAATCATAT ACTIGICITT TGTAAATAGC AGCITTIGIG TCATICICCC CACITTATTA	1260
35	GTTAATTTAA ATTOGAAAAA ACCCTCAAAC TAATATTCTT GTCTGTTCCA GTCTTATAAA	1320
	TAAAACTTAT AATGCATGTA AAAAAAAAAA A	1351
40		
70	(2)	
	(2) INFORMATION FOR SEQ ID NO: 105:	
45	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 2066 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 105:	
	GGCACGAGGC GGCGGAGGGC CACAATCACA GCTCCGGGCA TTGGGGGAAC CCGAGCCGGC	60
55	TGCGCCGGGG GAATCCGTGC GGGCGCCTTC CGTCCCGGTC CCATCCTCGC CGCGCTCCAG	120
55	CACCTCTGAA GTTTTGCAGC GCCCAGAAAG GAGGCGAGGA AGGAGGGAGT GTGTGAGAGG	180
	AGGGAGCAAA AAGCTCACCC TAAAACATTT ATTTCAAGGA GAAAAGAAAA	240
60	CAAAAATGGC TGGGGCAATT ATAGAAAACA TGAGCACCAA GAAGCTGTGC ATTGTTGCTGC	240

	GGATTCTGCT CGTGTTCCAA ATCATCGCCT TTCTGGTGGG AGGCTTGATT GCTCCAGGGC	360
5	CCACAACGGC AGTGTCCTAC ATGTCGGTGA AATGTGTGGA TGCCCGTAAG AACCATCACA	420
	AGACAAAATG GTTCGTGCCT TGGGGACCCA ATCATTGTGA CAAGATCCGA GACATTGAAG	480
	AGGCAATTCC AAGGGAAATT GAAGCCAATG ACATCGTGTT TTCTGTTCAC ATTCCCCTCC	540
10	CCCACATGGA GATGAGTCCT TGGTTCCAAT TCATGCTGTT TATCCTGCAG CTGGACATTG	600
	CCTTCAAGCT AAACAACCAA ATCAGAGAAA ATGCAGAAGT CTCCATGGAC GTTTCCCTGG	660
	CTTACCGTGA TGACGCATTT GCTGAGTGGA CTGAAATGGC CCATGAAAGA GTACCACGGA	720
15	AACTCAAATG CACCTTCACA TCTCCCAAGA CTCCAGAGCA TGAGGGCCGT TACTATGAAT	780
	GTGATGTCCT TCCTTTCATG GAAATTGGGT CTGTGGCCCA TAAGTTTTAC CTTTTAAACA	840
20	TCCGGCTGCC TGTGAATGAG AAGAAGAAAA TCAATGTGGG AATTGGGGAG ATAAAGGATA	900
	TCCGGTTGGT GGGGATCCAC CAAAATGGAG GCTTCACCAA GGTGTGGTTT GCCATGAAGA	960
25	CCTTCCTTAC GCCCAGCATC TTCATCATTA TGGTGTGGTA TTGGAGGAGG ATCACCATGA	1020
25	TGTCCCGACC CCCAGTGCTT CTGGAAAAAG TCATCTTTGC CCTTGGGATT TCCATGACCT	1080
	TTATCAATAT CCCAGTGGAA TGGTTTTCCA TCGGGTTTGA CTGGACCTGG ATGCTGCTGT	1140
30	TIGGTGACAT CCGACAGGGC ATCTTCTATG CGATGCTTCT GTCCTTCTGG ATCATCTTCT	1200
	GTGGCGAGCA CATGATGGAT CAGCACGAGC GGAACCACAT TGCAGGGTAT TGGAAGCAAG	1260
25	TOGGACCCAT TGCCGTTGGC TCCTTCTGCC TCTTCATATT TGACATGTGT GAGAGAGGGG	1320
35	TACAACTCAC GAATCCCTTC TACAGTATCT GGACTACAGA CATTGGAACA GAGCTGGCCA	1380
	TOGECTTCAT CATCOTGGCT GGAATCTGCC TCTGCCTCTA CTTCCTGTTT CTATGCTTCA	1440
40	TOGTATTICA GGTGTTTCGG AACATCAGTG GGAAGCAGTC CAGCCTGCCA GCTATGAGCA	1500
	AAGTCCGGCG GCTACACTAT GAGGGGCTAA TTTTTAGGTT CAAGTTCCTC ATGCTTATCA	1560
45	CCTTGGCCTG CGCTGCCATG ACTGTCATCT TCTTCATCGT TAGTCAGGTA ACGGAAGGCC	1620
45	ATTOGAAATG GGGCGGCGTC ACAGTCCAAG TGAACAGTGC CTTTTTCACA GGCATCTATG	1680
	GGATGTGGAA TCTGTATGTC TTTGCTCTGA TGTTCTTGTA TGCACCATCC CATAAAAACT	1740
50	ATGGAGAAGA CCAGTCCAAT GGAATGCAAC TCCCATGTAA ATCGAGGGAA GATTGTGCTT	1800
	TGTTTGTTTC GGAACTTTAT CAAGAATTGT TCAGCGCTTC GAAATATTCC TTCATCAATG	1860
	ACAACGCAGC TTCTGGTATT TGAGTCAACA AGGCAACACA TGTTTATCAG CTTTGCATTT	1920
55	GCAGTIGICA CAGTCACATT GATTGTACTT GTATACGCAC ACAAATACAC TCATTTAGCC	1980
	TTTATCTCAA AATGITAAAT ATAAGGAAAA AAGCGTCAAC AATAAATATT CTTGAGTATA	2040
60	AAAAAA AAAAAAAA AAAAAA	2066

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5 (2) INFORMATION FOR SEQ ID NO: 106:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1705 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:

15	AATTCOGCAK AGGGCAGCTG TCGGCTGGAA GGAACTGGTC TGCTCACACT TGCTGGCTTG	60
	CGCATCAGGA CTGGCTTTAT CTCCTGACTC ACGGTGCAAA GGTGCACTCT GCGAACGTTA	120
00	AGTCCGTCCC CAGCGCTTGG AATCCTACGG CCCCCACAGC CGGATCCCCT CAGCCTTCCA	180
20	GGTCCTCAAC TCCCGYGGAC GCTGAACAAT GGCCTCCATG GGGCTACAGG TAATGGGCAT	240
	CGCGCTGGCC GTCCTGGGCT GGCTGGCCGT CATGCTGTGC TGCGCGCTGC CCATGTGGCG	300
25	CGTGACGGCC TTCATCGGCA GCAACATTGT CACCTCGCAG ACCATCTGGG AGGGCCTATG	360
	GATGAACTGC GTGGTGCAGA GCACCGGCCA GATGCAGTGC AAGGTGTACG ACTCGCTGCT	420
	GGCACTGCCG CAGGACCTGC AGGCGGCCCG CGCCCTCGTC ATCATCAGCA TCATCGTGGC	480
30	TGCTCTGGGC GTGCTGCTGT CCGTGGTGGG GGGCAAGTGT ACCAACTGCC TGGAGGATGA	540
	AAGCGCCAAG GCCAAGACCA TGATCGTGGC GGGCGTGGTG TTCCTGTTGG CCGGCCTTAT	600
35	GGTGATAGTG CCGGTGTCCT GGACGGCCCA CAACATCATC CAAGACTTCT ACAATCCGCT	660
	GGTGGCCTCC GGGCAGAAGC GGGAGATGGG TGCCTCGCTC TACGTCGGCT GGGCCGCCTC	720
	CGGNCTGCTG CTCCTTGGCG GGGGGCTGCT TTGCTGCAAC TGTCCACCCC GCACAGACAA	780
40	GCCTTACTCC GCCAAGTATT CTGCTGCCCG CTCTGCTGCT GCCAGCAACT ACGTGTAAGG	840
• -	TGCCACGGCT CCACTCTGTT CCTCTCTGCT TTGTTCTTCC CTGGACTGAG CTCAGCGCAG	900
45	GCTGTGACCC CAGGAGGGCC CTGCCACGGG CCACTGGCTG CTGGGGACTG GGGACTGGGC	960
	AGAGACTGAG CCAGGCAGGA AGGCAGCAGC CTTCAGCCTC TCTGGCCCAC TCGGACAACT	1020
	TCCCAAGGCC GCCTCCTGCT AGCAAGAACA GAGTCCACCC TCCTCTGGAT ATTGGGGAGG	1080
50	GACGGAAGTG ACAGGGTGTG GTGGTGGAGT GGGGAGCTGG CTTCTGCTGG CCAGGATGGC	1140
		1200
<i></i>	TTAACCCTGA CTTTGGGATC TGCCTGCATC GGTGTTGGCC ACTGTCCCCA TTTACATTTT	
55	CCCCACTCTG TCTGCCTGCA TCTCCTCTGT TGCGGGTAGG CCTTGATATC ACCTCTGGGA	1260
	CTGTGCCTTG CTCACCGAAA CCCGCGCCCA GGAGTATGGC TGAGGCCTTG CCCACCCACC	1320
60	TCCCTCCGAA GTCCAGACTC GATCGACCCC TTTAGACCCG AGGGCCGAAG GTCCTCTAAA	1380

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	CAGGTTTGGG	CACTGGTGGG	GGAGGGGGCC	AGAGAGGCGG	CTCAGGTTGC	CCAGCTCTGT	:	1440
	GGCCTCAGGA	CTCTCTGCCT	CACCCGCTTC	AGCCCAGGGC	CCCTGGAGAC	TGATCCCCTC	:	1500
5	TGAGTCCTCT	GCCCCTTCCA	AGGACACTAA	TGAGCCTGGG	AGGGTGGCAG	GGAGGAGGGG	:	1560
	ACAGCTTCAC	CCTTGGAAGT	CCTGGGGTTT	TTCCTCTTCC	TTCTTTGTGG	TTTCTGTTTT		1620
10	GTAATTTAAG	AAGAGCTATT	CATCACTGTA	ATTATTATTA	TTTTCTACAA	TAAATGGGAC		1680
	CTGTGCACAG	GRAAAAAAAA	AAAAG					1705

15

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(2) INFORMATION FOR SEQ ID NO: 107:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1167 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:

25 TGCAGGAATT CGGCAGAGGT TTTCCGCTAG ACTCTGGCAG TTGGTGAGCA TCATGGCAAC 60 CGTTACAGCC ACAACCAAAG TCCCGGAGAT CCGTGATGTA ACAAGGATTG AGCGAATCGG 120 TGCCCACTCC CACATCCGGG GACTGGGGCT GGACGATGCC TTGGAGCCTC GGCAGGCTTC 180 30 GCAAGGCATG GTGGGTCAGC TGGCGGCACG GCGGGGGGT GGCGTGGTGC TGGAGATGAT 240 CCGGGAAGGG AAGATTGCCG GTCGGGCAGT CCTTATTGCT GGCCAGCCGG GCACGGGGAA 300 35 GACGGCCATC GCCATGGGCA TGGCGCAGGC CCTGGGCCCT GACACGCCAT TCACAGCCAT 360 CGCCGGCAGT GAAATCTTCT CCCTGGAGAT GAGCAAGACC GAGGCGCTGA CGCAGGCCTT 420 480 CCGGCGGTCC ATCGGCGTTC GCATCAAGGA GGAGACGGAG ATCATCGAAG GGGAGGTGGT 40 GGAGATCCAG ATTGATCGAC CAGCAACAGG GACGGGCTCC AAGGTGGGCA AACTGACCCT 540 CAAGACCACA GAGATGGAGA CCATCTACGA CCTGGGCACC AAGATGATTG AKTCCCTGAC 600 45 CAAGGACAAG GTCCAGGCCG GGGACGTGAT CACCATCGAC AAGGCGACGG GCAAGATCTC 660 CAAGCTGGGC CGCTCCTTCA CACGCGCCCG CGAACTACGA CGCTATGGGC TCCCAGACCA 720 AGTTCGTGCA GTGCCCAGAT GGGGAGCTCC AGAAACGCAA GGAGGTGGTG CACACCGTGT 50 780 CCCTGCACGA GATCGACGTC ATCAACTCTC GCACCCAGGG CTTCCTGGCG CTCTTCTCAG 840 GTGACACAGG GGAGATCAAG TCAGAAGTCC GTGAGCAGAT CAATGCCAAG GTGGCTGAGT 900 55 GOCGCGAGGA GGGCAAGGCG GAGATCATCC CTGGAGTGCT GTTCATCGAC GAGGTCCACA 960 TECTEGACAT CGAGAGCTTC TCCTTCCTCA ACCGGCCCT GGAGAGTGAC ATGGCGCCTG 1020 1080 TOCAGCAGGT CTATGGGGAT GCCGTGAGGG CTCTGGTAGC TGGTGCCCCG GATTCGCGTG 60

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359

	ATGCCACGGT TGGTGGCCTC GTGCCGAATT CCTGCAGCCC GGGGGATCCA CTAGTTCTAG	1140
5	AGCGGCCGCC ACCGCGGTGG ANCTCCN	1167
10	(2) INFORMATION FOR SEQ ID NO: 108: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1907 base pairs (B) TYPE: nucleic acid	
15	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 108:	
20	GGCACAGGGG AATCATCGTG TGATGTGTGT GCTGCCTTTG TGAGTGTGTG GAGTCCTGCT	60
	CAGGTGTTAG GTACAGTGTG TTTGATCGTG GTGGCTTGAG GGGAACCCTT GTTCAGAGCT	120
	GTGACTGCGG CTGCACTCAG AGAAGCTGCC CTTGGCTGCT CGTAGCGCCG GGCCTTCTCT	180
25	CCTCGTCATC ATCCAGAGCA GCCAGTGTCC GGGAGGCAGA AGGTACCGGG GCAGCTACTG	240
	GAGGACTGTG CGGGCCTGCC TGGGCTGCCC CCTCCGCCGT GGGGCCCTGT TGCTGCTGTC	300
30	CATCTATTIC TACTACTCCC TCCCAAATGC GGTCGGCCCG CCCTTCACTT GGATGCTTGC	360
	CCTCCTGGGC CTCTCGCAGG CACTGAACAT CCTCCTGGGC CTCAAGGGCC TGGCCCCAGC	420
	TGAGATCTCT GCAGTGTGTG AAAAAGGGAA TTTCAACGTG GCCCATGGGC TGGCATGGTC	480
35	ATATTACATC GGATATCTGC GGCTGATCCT GCCAGAGCTC CAGGCCCGGA TTCGAACTTA	540
	CAATCAGCAT TACAACAACC TGCTACGGGG TGCAGTGAGC CAGCGGCTGT ATATTCTCCT	600
40	CCCATTGGAC TGTGGGGTGC CTGATAACCT GAGTATGGCT GACCCCAACA TTCGCTTCCT	660
	GGATAAACTG CCCCAGCAGA CCGGTGACCG TGCTGGCATC AAGGATCGGG TTTACAGCAA	720
	CAGCATCTAT GAGCTTCTGG AGAACGGGCA GCGGGGGGGC ACCTGTGTCC TGGAGTACGC	780
45	CACCCCCTTG CAGACTITGT TTGCCATGTC ACAATACAGT CAAGCTGGCT TTAGCGGGGA	840
	GGATAGOCTT GAGCAGOCCA AACTCTTCTG CCGGACACTT GAGGACATCC TGGCAGATGC	900
50	CCCTGAGTCT CAGAACAACT GCCGCCTCAT TGCCTACCAG GAACCTGCAG ATGACAGCAG	960
	CTICTCGCTG TCCCAGGAGG TTCTCCGGCA CCTGCGGCAG GAGGAAAAGG AAGAGGTTAC	1020
	TGTGGGCAGC TTGAAGACCT CAGCGGTGCC CAGTACCTCC ACGATGTCCC AAGAGCCTGA	1080
55	GCTCCTCATC AGTGGAATGG AAAAGCCCCT CCCTCTCCGC ACGGATTTCT CTTGAGACCC	1140

AGGGTCACCA GGCCAGAGCC TCCAGTGGTC TCCAAGCCTC TGGACTGGGG GCTCTCTTCA

CTCCCTGAAT GTCCAGCAGA GCTATTTCCT TCCACAGGGG GCCTTGCAGG GAAGGGTCCA

60

1200

360

•	GGACTTGACA	TCTTAAGATG	CGTCTTGTCC	CCTTGGGCCA	GTCATTTCCC	CTCTCTGAGC	1320
	CTCGGTGTCT	TCAACCTGTG	AAATGGGATC	ATAATCACTG	CCTTACCTCC	CTCACGGTTG	1380
5	TTGTGAGGAC	TGAGTGTGTG	GAAGTTTTTC	ATAAACTTTG	GATGCTAGTG	TACTTAGGGG	1440
	GTGTGCCAGG	TGTCTTTCAT	GGGCCTTCC	AGACCCACTC	CCCACCCTTC	TCCCCTTCCT	1500
10	TTGCCCGGGG	ACGCCGAACT	CTCTCAATGG	TATCAACAGG	CTCCTTCGCC	CTCTGGCTCC	·1560
10	TOGTCATGTT	CCATTATTGG	GGAGCCCCAG	CAGAAGAATG	GAGAGGAGGA	GGAGGCTGAG	1620
	TTTGGGGTAT	TGAATCCCCC	GGCTCCCACC	CTGCAGCATC	AAGGTTGCTA	TGGACTCTCC	1680
15	TGCCGGGCAA	CTCTTGCGTA	ATCATGACTA	TCTCTAGGAT	TCTGGCACCA	CTTCCTTCCC	1740
	TOGCCCCTTA	AGCCTAGCTG	TGTATCGGCA	CCCCCACCCC	ACTAGAGTAC	TCCCTCTCAC	1800
20	TIGCGGTTTC	CTTATACTCC	ACCCCTTTCT	CAACGGTCCT	TTTTTAAAGC	ACATCTCAGA	1860
	TTAAAAAAAA	ААААААААА	ААААААААА	AAAAAAAGGG	CGGCCGC		1907

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(2) INFORMATION FOR SEQ ID NO: 109:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 611 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 109:

ATGAATTAAC GCCAAGCTNT NAATAGGGAC TCACTATGGG GGAAAGNIGG GTAACGCCTG 60 CAGGTACCGT TCCGGAATTC CCGGGTCGAC CCACGCGTCC GATGGGGCTT TAGTAAATCA 120 GGCTTGCAGG CTCAAAGCTG CAATCTGCCC ACTCTCAGGT ACTGAGACTT TGTGGGCCTC 180 AGACACCAGG AAGAAAGTTG GGATACAGTC ATTTGAGTTA AAAAGGGAAT GACCCCTCAG 240 AAACCCGCAT TAGCAGTGTT ACTCTTGGAA GTGCCTTTAC TTTTAACGCT CTCTGTTCTG 300 AAAAAGAGGT GTTTGGTTAC GTGTGAGCCA ACATCACGTT TTGTTAGCTG TGATTTACCT 360 TTGTCCGTTT AAAAGACTTC ACGGAGCCAT TCTGTATACA AGGTGTGCTC TTTCCAATGT 420 AGAAGGGGTT ATGGAAAAGG GTGCGATCCT TTGCTGTAAA CTGGAGAGAC CAGTCCCAAA 480 CAGAGGGGAA TTTTAAGCCC TTCTCATCAC CCAATTGGAT GTTTTTGCTT ATAGCAAATT 540 ССТССАВАЯТ АВАТАВАТАВ АТАТТТССАВ ВАСТАВАВАВ ВАВАВАВАВА ВАВАВАВАВА 600 GGGGGGNCCN C 611

361

(2) INFORMATION FOR SEQ ID NO: 110:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2632 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:

10 TCCCAGCTCT CAGGACAAGG GCCCTGGGCG ATCTTTTAAA AAAGCCGATT GGGTGTCTTT 60 CTAAAANTAC AACCAGTACT TCATCGTCAA GTTTCTGGGA AGGGAGTCCC CTCCAGATTC 120 15 TCATGGAGTG ACAAATCTTG ACTCTTGCTC CTGGAATTTT TCAGGCCCAA ACTAGCGTTT 180 CTACAATGAT TTATTTGGCA AATTTGTCTT GATTATGGGT GGCTGATGAG GAACGTGCTT 240 TTGTTAGGAA CCGAAACTGG GCGCGGTGA GGGCGTGTAC GCAATGAGTC CGGAAGAGGG 300 20 TGAAATGCTT TCGGTAGGCA CTCCACGGCT GTGAAGATGG CGGCGGCTGC GTGGCTTCAG 360 GTGTTGCCTG TCATTCTTCT GCTTCTGGGA GCTCACCCGT CACCACTGTC GTTTTTCAGT 25 GCGGGACCGG CAACCGTAGC TGCTGCCGAC CGGTCCAAAT GGCACATTCC GATACCGTCG 480 GGGAAAAATT ATTITAGTTT TGGAAAGATC CTCTTCAGAA ATACCACTAT CTTCCTGAAG 540 TTTGATGGAG AACCTTGTGA CCTGTCTTTG AATATAACCT GGTATCTGAA AAGCGCTGAT 600 30 TGTTACAATG AAATCTATAA CTTCAAGGCA GAAGAAGTAG AGTTGTATTT GGAAAAACTT 660 AAGGAAAAAA GAGGCTTGTC TGGGAAATAT CAAACATCAT CAAAATTGTT CCAGAACTGC 720 35 AGTGAACTCT TTAAAACACA GACCTTTTCT GGAGATTTTA TGCATCGACT GCCTCTTTTA 780 GGAGAAAAAC AGGAGGCTAA GGAGAATGGA ACAAACCTTA CCTTTATTGG AGACAAAACC 840 GCAATGCATG AACCATTGCA AACTTGGCAA GATGCACCAT ACATTTTTAT TGTACATATT 900 40 GGCATTTCAT CCTCAAAGGA ATCATCAAAA GAAAATTCAC TGAGTAATCT TTTTACCATG 960 ACTOTTGAAG TGAAGGGTCC CTATGAATAC CTCACACTTG AAGACTATCC CTTGATGATT 1020 45 TTTTCATGG TGATGTGTAT TGTATATGTC CTGTTTGGTG TTCTGTGGCT GGCATGGTCT 1080 GCCTGCTACT GGAGAGATCT CCTGAGAATT CAGTTTTGGA TTGGTGCTGT CATCTTCCTG GGAATGCTTG AGAAAGCTGT CTTCTATGCG GAATTTCAGA ATATCCGATA CAAAGGARAA 1200 50 TCTGTCCAGG GTGCTTTGAT CCTTGCAGAR CTGCTTTCAG CAGTGAAACG CTCACTGGCT 1260 CGAACCCTGG TCATCATAGT CAGTCTGGGA TATGGCATCG TCAAGCCACG CCTGGAGTCA 1320 55 CTCTTCATAA GGTTGTAGTA GCAGRAGCCC TCTATCTTTT GTTCTCTGGC ATGGAAGGGG 1380 TCCTCAGAGT TACTGGGGCC CAGACTGATC TTGCTTCCTT GGCCTTTATC CCCTTGGCTT 1440 TCCTAGACAC TGCCTTGTGC TGGTGGATAT TTATTAGCCT GACTCAAACA ATGAAGCTAT 1500 60

240

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•	TAAAACTICG GAGGAACATT GIAAAACTCT CITIGIATCG GCATTICACC AACACGCTTA	1560
	TTTTGGCAGT GGCAGCATCC ATTGTGTTTA TCATCTGGAC AACCATGAAG TTCAGAATAG	1620
5	TGACATGTCA GTCGGACTGG CGGGAGCTGT GGGTAGACGA TGCCATCTGG CGCTTGCTGT	1680
	TCTCCATGAT CCTCTTTGTC ATCATGGTTC TCTGGCGACC ATCTGCAAAC AACCAGAGGT	1740
10	TTGCCTTTTC ACCATTGTCT GAGGAAGAGG AGGAGGATGA ACAAAAGGAG CCTATGCTGA	1800
10	AAGAAAGCTT TGAAGGAATG AAAATGAGAA GTACCAAACA AGAACCCAAT GGAAATAGTA	1860
	AAGTTAACAA AGCACAGGAA GATGATTTGA AGTGGGTAGA AGAGAATGTT CCTTCTTCTG	1920
15	TGACAGATGT AGCACTTCCA GCCCTTCTGG ATTCAGATGA GGAACGAATG ATCACACACT	1980
	TTGAAAGGTC CAAAATGGAG TAAGGAATGG GAAGATTTGC AGTTAAAGAT GGCTACCATC	2040
20	AGGGAAGAGA TCAGCATCTG TGTCAGTCTT CTGTACGGCT CCATGGGATT AAAGGAAGCA	2100
20	ATGACATCCT GATCTGTTCC TTGATCTTTG GGCATTGGAG TTGGCGAGAG GTGTCAGAAC	2160
	AAAGAGAACA TCTTACTGAA AACAAGTTCA TAAGATGAGA AAAATCTACG AGCTTCTTAT	2220
25	TTACAACACT GCTGCCCCCT TTCCTCCCAG ACTCTGACAT GGATGTTCAT GCAACTTAAG	2280
	TGTGTTGTTC CTGAACTTTC TGTAATGTTT CATTTTTTAA ATCTGACAAA CTAAAAAGTT	2340
30	TAACGICTIC TAAAAGATIG TCATCAACAC CATAATATGT AATCICCAGG AGCAACIGCC	2400
50	TGTAATTTTT ATTTATTTAG GGAGTTACAT AGGTGATGGG GGAAATTGTT AACTACCTTT	2460
	CATTITICCTG GGAAGTCAAG GTTACATCTT GCAGAGGTTG TTTTGAGAAA AAAGGGCCCT	2520
35	TCTGAGTTAA GGAGCCATAG TTCTATCAAT GATCAAAAGA AAAAAAAAA AACTCGATCG	2580
	GCACGAGGG GGGCCCGGTA CCCAATTCGC CCTATGGGAN TCGAATGAGA CC	2632
40		
	(2) INFORMATION FOR SEQ ID NO: 111:	
•	• • • • • • • • • • • • • • • • • • •	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2249 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 111:	
	GAATTOGGCA CGAGCTCACC GTGCTGCGTG ACACAAGGCC AGCCTGCGCC TACGAGCCCA	60
	TOGACTITIKT RATOGCCCTC ATCTACGACA TOGTACTGSW TOTGGTCACC CTGGGGCTGG	120
55	CONTROLLE MONOTOCCE A ACTIONA ACA COMPOSA ACOM CA ACCOCCO MINISTRAÇÃO	18

TCACAGCCTT CCTCTCTGTG CTCATCTGGG TGGCCTGGAT GACCATGTAC CTCTTCGGCA

60 ATGTCAAGCT GCAGCAGGG GATGCCTGGA ACGACCCCAC CTTGGCCATC ACGCTGGCGG

	CCAGCGCTGG GTCTTCGTCA TCTTCCACGC CATCCCTGAG ATCCACTGCA CCCTTCTGCC	360
5	AGCCCTGCAG GAGAACACGC CCAACTACTT CGACACGTCG CAGCCCAGGA TGCGGGAGAC	420
J	GCCCTTCGAG GAGGACGTGC AGCTGCCGCG GGCCTATATG GAGAACAAGG CCTTCTCCAT	480
	GGATGAACAC AATGCAGCTC TCCGAACAGC AGGATTTCCC AACGGCAGCT TGGGAAAAAG	540
10	ACCCAGTGGC AGCTTGGGGA AAAGACCCAG CGCTCCGTTT AGAAGCAACG TGTATCAGCC	600
	AACTGAGATG GCCGTCGTGC TCAACGGTGG GACCATCCCA ACTGCTCCGC CAAGTCACAC	660
15	AGGAAGAMAC CITTGGTGAA AGACTITAAG TTCCAGAGAA TCAGAATTTC TCTTACCGAT	720
13	TIGCCTCCCT GCCTGTGTCT TTCTTGAGGG AGAAATCGGT AACAGTTGCC GAACCAGGCC	780
	GCCTCACAGC CAGGAAATTT GGAAATCCTA GCCAAGGGGA TTTCGTGTAA ATGTGAACAC	840
20	TGACGAACTG AAAAGCTAAC ACCGACTGCC CGCCCCTCCC CTGCCACACA CACAGACACG	900
	TAATACCAGA CCAACCTCAA TCCCCGCAAA CTAAAGCAAA GCTAATTGCA AATAGTATTA	960
25	GGCTCACTGG AAAATGTGGC TGGGAAGACT GTTTCATCCT CTGGGGGTAG AACAGAACCA	1020
23	AATTCACAGC TGGTGGGCCA GACTGGTGTT GGTTGGAGGT GGGGGGCTCC CACTCTTATC	1080
	ACCTCTCCCC AGCAAGTGCT GGACCCCAGG TAGCCTCTTG GAGATGACCG TTGCGTTGAG	1140
30	GACAAATGGG GACTTTGCCA CCGGCTTTGC CTGGTGGTTT GCACATTTCA GGGGGTCAG	1200
	GAGAGITAAG GAGGITGIGG GIGGGATICC AAGGIGAGGC CCAACIGAAT CGIGGGGIGA	1260
35	GCTTTATAGC CAGTAGAGGT GGAGGGACCC TGGCATGTGC CAAAGAAGAG GCCCTCTGGG	1320
JJ	TGATGAAGTG ACCATCACAT TTGGAAAGTG ATCAACCACT GTTCCTTCTA TGGGGCTCTT	1380
	SCTCTAGTGT CTATGGTGAG AACACAGGCC CCGCCCCTTC CCTTGTAGAG CCATAGAAAT	1440
40	ATTCTGGCTT GGGGCAGCAG TCCCTTCTTC CCTTGATCAT CTCGCCCTGT TCCTACACTT	1500
	ACGGGTGTAT CTCCAAATCC TCTCCCAATT TTATTCCCTT ATTCATTTCA AGAGCTCCAA	1560
45	TOGGGTCTCC AGCTGAAANS CCCTCCGGGA GGCAGGTTGG AAGGCAGGCA CCACGGCAGG	1620
43	TTTTCCGCGA TGATGTCACC TAGCAGGGCT TCAGGGGTTC CCACTAGGAT GCAGAGATGA	1680
	CCTCTCGCTG CCTCACAAGC AGTGACACCT CGGGTCCTTT CCGTTGCTAT GGTGAAAATT	1740
50	CCTGGATGGA ATGGATCACA TGAGGGTTTC TTGTTGCTTT TGGAGGGTGT GGGGGATATT	1800
	TTGTTTTGGT TTTTCTGCAG GTTCCATGAA AACAGCCCTT TTCCAAGCCC ATTGTTTCTG	1860
55	TCATGGTTTC CATCTGTCCT GAGCAAGTCA TTCCTTTGTT ATTTAGCATT TCGAACATCT	1920
55	CGGCCATTCA AAGCCCCCAT GITCTCTGCA CTGTTTGGCC AGCATAACCT CTAGCATCGA	1980
	TTCAAAGCAG AGTTTTAACC TGACGGCATG GAATGTATAA ATGAGGGTGG GTCCTTCTGC	2040
60	AGATACTCTA ATCACTACAT TOCTTTTTCT ATAAAACTAC CCATAAGCCT TTAACCTTTA	2100

364

	AAGAAAAATG AAAAAGGTTA GTGTTYGGGG GCCGGGGGAG GACTGACCGC TTCATAAGCC	2160
5	AGTACGICIG AGCTEAGIAT GITTCAAFAA ACCTITIGAT ATTICICAAA AAAAAAAAA	2220
•	AAAAANCCCG GGGGGGGGCC CGGACCTGG	2249
10	(2) DESCRIPTION FOR CES = 10. 112.	
	(2) INFORMATION FOR SEQ ID NO: 112:	
	(i) SEQUENCE CHAPACTERISTICS: (A) LENGTH: 2193 base pairs	
15	(3) TYPE: mucleic acid	
	(C) STRAUTENESS: double	
	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 112:	
	GATACTATAA GGCAAGTGAC TCACGGGTGC GCCGTTAGAC TAGTGGATCC CGGGTGCAGG	60
	AATTOGGCAG AGCGCCGCCG GAGCCGAAGT GCTGGCGCCCC CCGCGGCCGC TGCCTCCGCG	120
25	GANCOCAAAA TCATGAAAST CACCSTGAAG ACCOCGAAGA AAAGGAGGAA TTCGCCGTGC	180
	CCGAGAATAG CTCCGTCCAG CAGTTTAAGG AAGAAATCTC TAAACGTTTT AAATCACATA	240
30	CTGACCAACT TGTGTTGAJA TTTGCTGGAA AAATTTTGAA AGATCAAGAT ACCTTGAGTC	300
•	AGCATGGAAT TCATGATGGA CTTACTGTTC ACCTTGTCAT TAAAACACAA AACAGGCCTC	360
	AGGATCATTC AGCTCAGCAA ACAAATACAG CTGGAAGCAA TGTTACTACA TCATCAACTC	420
35	CTAATAGTAA CTCTACATCT GGTTCTGCTA CTAGCAACCC TTTTGGTTTA GGTGGCCTTG	480
	GGGGACTTGC AGGTCTGAGT AGCCTGGGTT TGAATACTAC CAACTTCTCT GAACTACAGA	540
40	GTCAGATGCA GCGACAACTT TTGTCTAACC CTGAAATGAT GGTCCAGATC ATGGAAAAWC	600
	CCYTTGTTCA GAGCATGCTC MTCAAATCCT GACCTGATGN AGACAGTTAA TTATGGCCAA	660
	TCCACAAATG CAGCAGTTGA TACAGAGAAA TCCCAGAAAT TAGTCATATG TTGAATAATC	720
45	CAGATATAAT GAGACAAAGG TTGGAACTTG CCCAGGAATC CAGCAATGAT GCAGGAGATG	780
	ATGAGGAACC AGGACCGAGC TITGAGCAAC CTAGAAAGCA TCCCAGGGGG ATATAATGCT	840
50	TTAAGGCGCA TGTACACAGA TATTCAGGAA CCAATGCTGA GTGCTGCACA AGAGCAGTTT	900
-	GGTGGTAATC CATTTGCTC CTTGGTGAGC AATACATCCT CTGGTGAAGG TAGTCAACCT	960
	TCCCGTACAG AAAATAGAGA TCCACTACCC AATCCATGGG CTCCACAGAC TTCCCAGAGT	1020
55	TCATCAGCTT CCAGCGGCAC TGCCAGCACT GTGGGTGGCA CTACTGGTAG TACTGCCAGT	1080
	GGCACTTCTG GGCAGAGTAC TACTGCGCCA AATTTGGTGC CTGGAGTAGG AGCTAGTATG	1140

TTCAACACAC CAGGAATECA GAGETTGTTG CAACAAATAA CTGAAAACCC ACAACTTATG

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•	CAAAACATGT TGTCTGCCCC CTACATGAGA AGCATGATGC AGTCACTAAG ÇCAGAATCCT	1260
	GACCTTGCTG CACAGATGAT GCTGAATAAT CCCCTATTTG CTGGAAATCC TCAGCTTCAA	1320
5	GAACAAATGA GACAACAGCT CCCAACTTTC CTCCAACAAA TGCAGAATCC TGATACACTA	1380
	TCAGCAATGT CAAACCCTAG AGCAATGCAG GCCTTGTTAC AGATTCAGCA GGGTTTACAG	1440
10	ACATTAGCAA CGGAAGCCCC GGGCCTCATC CCAGGGTTTA CTCCTGGCTT GGGGGCATTA	1500
10	GGAAGCACTG GAGGCTCTTC GGGAACTAAT GGATCTAACG CCACACCTAG TGAAAACACA	1560
	AGTCCCACAG CAGGAACCAC TGAACCTGGA CATCAGCAGT TTATTCAGCA GATGCTGCAG	1620
15	GCTCTTGCTG GAGTAAATCC TCAGCTACAG AATCCAGAAG TCAGATTTCA GCAACAACTG	1680
	GAACAACTCA GTGCAATGGG ATTTTTGAAC CGTGAAGCAA ACTTGCAAGC TCTAATAGCA	1740
20	ACAGGAGGTG ATATCAATGC AGCTATTGAA AGGTTACTGG GCTCCCAGCC ATCATAGCAG	1800
20	CATTICTGTA TCTKGAAAAA ATGTAATTTA TTTTTGATAA CGGCTCTTAA ACTTTAAAAT	1860
	ACCIGCITIA TITCATITIG ACTOTIGGAA TICTGIGCIG TIATAAACAA ACCCAATATG	1920
25	ATGCATTITA AGGTGGAGTA CAGTAAGATG TGTGGGTTTT TCTGTATTIT TCTTTTCTGG	1980
	AACAGTGGGA ATTAAGGCTA CTGCATGCAT CACTTCTGCA TTTATTGTAA TTTTTTAAAA	2040
30	ACATCACCIT TTATAGTTGG GIGACCAGAT TTTGTCCTGC ATCTGTCCAG TTTATTTGCT	2100
	TTTTAAACAT TAGCCTATGG TAGTAATTTA TGTAGAATAA AAGCATTAAA AAGAAGCAAA	2160
	AAAAAAAAA AAAAATTCCT GCGCCCGCGA ATTCTTCT	2198
35		
	(2) INFORMATION FOR SEQ ID NO: 113:	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1043 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
45	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 113:	
	CTGAAGTGTA TGTGGTGAGG AAGAAGAGGC TCCTACTGTA GACAGCCTTG TTCTACAGAT	. 60
50	CCTCCCAGAA ATCTCTGGGC CAGGTGGAAC CCAGGGTCAG AGAGGGATGG GAGAGAGGTT	120
	TAATTTTCCA TGATAAATAA AAATCTATAA AATAATAAAC AAGAGAAAAG AGATTGGAAA	180
55	CAGCCAGGIT GGAGCAGIGA GIGAGIAAGG AAACCIGGCI GCCCICICCA GATICCCCAG	240
	GCTCTCAGAG AAGATCAGCA GAAAGTCTGC AAGACCCTAA GAACCATCAG CCCTCAGCTG	300
	CACCTCCTCC CCTCCAAGGA TGACAAAGGC GCTACTCATC TATTTGGTCA GCAGCTTTCT	360

60 тесссталат саевссаесс телтелетов стетелетте есселенос теслестева

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	RGACTTGGAT GGGTTTGAGG GTTACTCCCT GAGTGACTGG CTGTGCCTGG CTTTTGTGGA	480
5	AAGCAAGTTC AACATATCAA AGATWAATGA AAATGCAGAT GGAAGCTTTG ACTATGGSCT	540
3	CTTCCAGATC AACAGCCACT ACTGGTGCAA CRATTATAAG AGTTACTCGG AAAACCTTTG	600
	CCACGTAGAC TGTCAAGATC TGCTGAATCC CAACCTTCTT GCAGGCATCC ACTGCGCAAA	. 660
10	AAGGATTGTG TCCGGAGCAC GGGGGATGAA CAACTGGGTT AGAATGGAAG KTTGCACTGT	720
	TCAGGCCGGC CACTCTTCTA CTGGCTGACA GGATGCCGCC TGAGATKAAA CARGGTGCGG	780
15	GTGCACCGTG GARTCATTCC AAGACTCCTG TCCTCACTCA RGGATTCTTC ATTTCTTCTT	840
13	CCTACTGCCT CCACTTCATG TTATTTTCTT CCCTTCCCAT TTACAACTAA AACTGACCAG	900
	AGCCCCAGGA ATAAATGGTT TTCTTGGCTT CCTCCTTACT CCCATCTGGA CCCAGTCCCC	960
20	TOGTTCCTGT CTGTTATTTG TAAACTGAGG ACCACAATAA AGAAATCTTT ATATTTATCG	1020
	AAAAAAAAA AAAAAAAACT CGA	1043
25		
	(2) INFORMATION FOR SEQ ID NO: 114:	
30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 703 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:	
	GAATTCGGCA CGAGTGCGCG GGCACCACGG CGGTTTTTCG ACGCTGGCGG TGGACGCAGG	60
40	CAGCATGGAC CACGGTTGCT GGGCGGATGG GGAGCGTCTA TGGTCAGTTG CCTTAGAAGT	120
40	COTGAGATCG GAACCTCCAG TTCGAAGACC CTCGACGATC CCTGACAAGG GGATCTCTGA	186
	CACATGATTG GAGCTCTTTT TGAAATGTTT CTTGCCCTTC CTGGAGCAGA GGAGCCATTA	246
45	TTTATGCAGG TACATCGAAG TCTTTTGACC TCCATACAGT GATTATGCTT GTCATCGCTG	300
	GTGGTATCCT GGCGGCCTTG CTCCTGCTGA TAGTTGTCGT GCTCTGTCTT TACTTCAAAA	360
50	TACACAACGC GCTAAAAGCT GCAAAGGAAC CTGAAGCTGT GGCTGTAAAA AATCACAACC	420
50	CAGACAAGGT GTGGTGGGCC AAGAACAGCC AGGCCAAAAC CATTGCCACG GAGTCTTGTC	48
	CTGCCCTGCA GTGCTGTGAA GGATATAGAA TGTGTGCCAG TTTTGATTCC CTGCCACCTT	54
55	GCTGTTGCGA CATAAATGAG GGCCTCTGAG TTAGGAAAGG TGGGCACAAA AATCTTCATG	60

AGCAATACTT CTTAGTAGAT TGTTTTGTTA TTCAAATCAA GTTCTAGTGT TTTTATGTGA

GATTATATAA TTTACAGTGT TGTTTTATAT ACTTTTGAAT AAA

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(2)	INFORMATION	FOR	SEQ	ID	NO:	115:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3684 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:

GCCAGAGGGG GCATGAGCAG GAGGAGGATT ACCGCTACGA GGTGCTCACG GCCGAGCAGA 60 15 TTCTACAACA CATGGTGGNA ATGTATCCGG GAGGTCAACG AGGTCATCCA GAATCCAGCA 120 ACTATCACAA GAATACTCCT TAGCCACTTC AATTGGGATA AAGAGAAGCT AATGGAAAGG 180 20 TACTTTGATG GAAACCTGGA GAAGCTCTTT GCTGAGTGTC ATGTAATTAA TCCAAGTAAA 240 AAGTCTCGAA CACGCCAGAT GAATACAAGG TCATCAGCAC AGGATATGCC TTGTCAGATC 300 TGCTACTTGA ACTACCCTAA CTCGTATTTC ACTGGCCTTG AATGTGGACA TAAGTTTTGT 360 25 ATGCAGTGCT GGAGTGAATA TTTAACTACC AAAATAATGG AAGAAGGCAT GGGTCAGACT 420 ATTTCGTGTC CTGCTCATGG TTGTGATATC TTAGTGGATG ACAACACAGT TATGCGCCTG 30 ATCACAGATT CAAAAGTTAA ATTAAAGTAT CAGCATTTAA TAACAAATAG CTTTGTAGAG 540 TGCAATCGAC TGTTAAAGTG GTGTCCTGCC CCAGATTGCC ACCATGTTGT TAAAGTCCAA 600 TATCCTGATG CTAAACCTGT TCGCTGCAAA TGTGGGCGCC AATTTTGCTT TAACTGTGGA 660 35 GAAAATTGGC ATGATCCTGT TAAATGTAAG TGGTTAAAGA AATGGATTAA AAAGTGTGAT 720 GATGACAGTG AAACCTCCAA TTGGATTGCA GCCAACACAA AGGAATGTCC CAAATGCCAT 780 40 GTCACAATTG AGAAGGATGG TGGTTGTAAT CACATGGTCT GTCGTAACCA GAATTGTAAA 840 GCAGAGTTTT GCTGGGTGTG TCTTGGCCCA TGGGAACCAC ATGGATCTGC CTGGTACAAC 900 TGTAACCGCT ATAATGAGGA TGATGCAAAG GCAGCAAGAG ATGCACAGGA GCGATCTAGG 960 45 GCAGCCCTGC AGAGGTACCT GTTCTACTGT AATCGCTATA TGAACCACAT GCAGAGCCTG 1020 CGCTTTGAGC ACAAACTATA TGCTCAGGTG AAACAGAAAA TGGAGGAGAT GCAGCAGCAC 1080 50 AACATGTCCT GGATTGAGGT GCAGTTCCTG AAGAAGGCAG TTGATGTCCT CTGCCAGTGT 1140 CGTGCCACAC TCATGTACAC TTATGTCTTC GCTTTCTACC TCAAAAAGAA TAACCAGTCC 1200 1260 ATTATCTTTG AGAATAACCA AGCAGATCTA GAGAATGCCA CAGAGGTGCT CTCGGGCTAC 55 1320 TACAGATACT GTGAGAGTCG ACGAAGGGTT TTGTTACAGC ATGTGCATGA AGGCTATGAA 1380 60 AAAGATCTGT GGGAGTACAT TGAGGACTGA GAATGGCCCT GCATAAAATG AACTCTGAAA 1440

	ACTITACCAT	CTAGAGTGCT	CATGCAATTA	AAACAAAACA	AACACAAACA	AGGAGGCACT	1500
5	AAGCCTATTC	TGACACCACT	GGTCTGTAGT	ACCAGAATTG	TTTTGTTAAT	GGAAAGTTTA	1560
J	AGTAAATTAT	ATTGTAATAA	AAAGGTAGAT	AAACCATTGT	ACAACAGTAT	TCTAGGCCGC	1620
	CAACAAAAGT	GTGACAGACA	CACTAAAAGC	CCTCCAACTT	TAACTTGTAA	CGTAGCTTCA .	1680
10	TTCTCAAAGC	TGACTCCTTT	TTTTTCTTTT	TCCTTTTCCT	GAGTGTAGTA	CAGTTAAAAT	1740
	TTCAAACAGC	TCCTTGACAC	TGCTTTTCAT	GTTCAAACCA	GCCATTTTGT	TGTACTTTGG	1800
15	TAAAGGACCT	CTTCCCCTTC	CTCCCCTACA	CATACAGATA	CACCCACACA	CAGACTGACT	1860
15	CTCTTTCTCT	CATACCCCAA	GGTCATGAGT	GAATGATGCT	TAGTTCCTTG	TAAAGAAAAT	1920
	CTTGGGATGG	GGAAAGGGGT	AGGCAGCAAG	AGGATTCAAC	AAACGAAAAA	CATAAAAACT	1980
20	TTGTATATGA	CTTTTAAAAC	AAGAGGACAA	CACAGTATTT	TTCAAAATTG	TATATAGCGC	2040
	ATATGCATGG	ACAAAGCAAG	CCTCCCACCT	GTTTGCATAA	TGTTTAATTA	СААААААТА	2100
25	TTTATTCTTT	AAAAATCTTC	AAGATTATGT	CTATTTGCTG	TGCATTTTCT	TTCAGTTTGC	2160
	TTATCTTTCC	CGGGTTGGGG	TTGGGATAAA	GETETETCGE	TTTAGCACCT	CTGGAAGACC	2220
	TATCTAGAGC	TCTTTCACTT	TCCTGAGGTT	ATTTTGCCCY	TTCTGGTGTT	GGTATGTCTG	2280
30	TTGCCGGCCA	TOGGCTNCAY	GCCTTGAATT	CCTCCTCTTG	ATCAGGGACA	AGGGAGGTCA	2340
	AGCTCTGACT	AATGCCATGA	CCTGATTAAG	GGGTACAGCA	GGGAGTTTTG	TTGCTACAGC	2400
35	TCATGAATTA	ACCTGTCCCA	ACCTAATCCC	CCTCCATGGC	ATCATGCCTC	TACCCAAGCC	2460
55	TTTGTGTGCC	CATGTTATGC	ACACAGCTGT	AGGCATTCTT	AAGTCCCCTG	TCGCATCCAG	2520
	TGGAAGCATT	TTAAAATTTC	TITTACITTT	TGGTTTTCCC	TTAATTGCTG	CTTTTCAGAT	2580
40	TTTAGTTATG	GCTCGTCTGC	TCACCCCTTC	TCTACATTAG	GGTGTCAAAG	AGAATGTTTT	2640
	GCTTTAAATA	TAAATAGCCA	TTCATTTAGT	CTCAGATTGT	GAATTTAAAA	TGGTGGATAC	2700
45	CGAAATTGCT	TGTGTGTGTT	GCTGTGGGTT	TGGTTTGAAG	GCAAACACCC	CTAGAACATG	2760
	ATATTCCCAT	CTAGTGCATT	TAAATAGAAA	TCACTGAGTT	TGCTGCTTTT	TTATTGTCAG	2820
	CAGATAGGAG	AATTAATAAT	GCATTTTAGC	TGTGATGTCC	ATTTTTATGA	AATTCCTACT	2880
50	AAGAGCTATG	TTAAAAGTAA	AGGATGGTGG	TOGTTGTATT	AACTATATAC	CTGTTTAGGC	2940
	CATTCTGGCT	GIGGIATITI	TCAATAGGTC	AGCATCTGTA	AATCTGTCAG	TTTTATACAG	3000
55	GAGTGCAGAG	TGAACTAGGC	AACTAGATTA	AGAGGTCTAA	ATATGAAATA	CCACTTGAGG	3066
<i>33</i>	CTGAGGACCT	CTTCGTCTTC	CTTTAAATGT	CTTTTGCCTA	GGGAGTGTTT	ACCATTTGTG	312
	AGGCAGCTTT	GICTGCTCTT	ACACTGTACA	TCCTATTACT	CCATTGGGAA	GTAGGTTCAC	318
60	TTTCCTCTGG	CCTTTTGCCT	AAGTTAGGCT	TTGCTGAATC	AACCCTACTT	TTCCTTTTAG	324

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	AAAAGGTTGT	TACAGGAGAT	TTACTGGCAA	CIGITCITIT	CCCATCAAAA	ATCAGTGAAT	3300
۲	GTTTGCTGAG	TATAAATGCT	GCTTCCTTAA	ACCACTTGTC	GCTTTAGGAT	CAACTTTACC	3360
J	TGTACCTTTT	CTCCTTTCCT	CCCTTGCCAC	CTCAGGTGCA	AATCTGAACT	CAGTGTCTGC	3420
	TICTTCCATT	TICTCGTCTC	TCTCCCCTCT	TCCCCCATTA	TCCATATGAC	ATTATTTTAC .	3480
10	TTCAAATGAC	AGCATCAATC	TTAAAAAGAT	ATACATTAAA	ACTAAGGAGT	TTTTTTAAAG	3540
	AAAGCCTGAA	TAAGITCCTT	TCCCTGGTAA	CTTTGAAAAG	CAGTCAGAGT	TGCTATATAG	3600
15	ATATATGTGG	CTCCTTTAAA	ATGCTTTGTG	TATGTGTGGT	GTTTAAAAAA	AAAAAAAA	3660
13	TTCGGGGGGG	GGCCCGGTNC	CCAT				3684

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(2) INFORMATION FOR SEQ ID NO: 116:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1965 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 116:

AAGAAAGGGT ATTAAAATTC TAGATCACAT ATGGACCCGG GAAGGTTTTT NACCCTCTGT 60 120 TAGTGACATC GAGTCTCCCA CTAGACAAAA TAGGTGGAAA AATCTCTCGA GGGCTCACAT TGTTTTGTCA TCTTCAGGAA AAACACCACC AGGCCATACC ACAGCCTGCC CAGTGAGGCG 180 GTCTTTGCCA ACAGCACCGG GATGCTGGTG GTGGCCTTTG GGCTGCTGGT GCTCTACATC 240 CTTCTGGCTT CATCTTGGAA GCGCCCAGAG CCGGGGATCC TGACCGACAG ACAGCCCCTG CTGCATGATG GGGAGTGAAG CAGCAGGAAG GGGCTCCCAA GAGCTCCTGG TGGTGCAGCC 360 TGTGCTCCCC TCAGAAGCTC TGCTCTTCCC AGGGCTCCCG GCTGGTTTCA GCAGGCGACT TTCTTCCAAT GCTGGGCCCA GACTTCTTGC CTGGGTGCTG GCCTGCCCTC TCCGGNCCGC 480 TTGCTGCCTG TCTGCTTTCC TTGGTGGYTT TGCTGGGTGC TGGGCCTGCC CTCTCCGGCC 600 GCTTGCTGCC TGTCTGCTTT CCTTGGTGGC TTTGCTGGGT GCTGGGCCTG CCTTCTCTGG CTGCTTGCTG CCTGTCTGCT TTCCTTGGTG GCTTTGGCTT CTGCACTCCT TGGCGTCASC 660 TCTCAGGTCC TCCATTCACA CGAGGTCCTC CTCGCTCTGG CCGCTCTTGC TGCTCCTGTC · 720 TGAAGAWATC AGACTGATTT CCTCTTAAGA CTCCTAGGGA TGTGGTGAAG AGCTGGGACT 780 CAACTCCAGT CCACCGTCTG AAACATGAGG GARGTGAGGT GTCCGTCCAC TTCCCCCATA 840 AAGGTGTGCA TTTCAGTTAG GCTGCCCCGC CACAGAGCAG GCTTCATCTG CTCTGCCATC 900

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	CAGCCCCATC TOGATGTGAG GTGGGGTGGA GACATCATGG GGTGATTGCA GAAAGGGGGA	960
	GTGGCGGCCC ACGCAGCTTC TGCTGAGGAG CTGACCGCTC TGAGCTGTTC TGTTTCGTAT	1020
5	TOCTOCTCTG TGTCTGCATG TATTGTGACC GTGCGGCTCC ACCTCTTCCA GCTGCTGCTA	1080
	CASCITGAGGC CIGGATCCCG GCCTTTCCCT GTGACTTACG TGTCTGTCAC CGGCANGCAG	1140
10	CCCTACAAAT CCTGGTGACC TGCTCTCCCA AGAACAGAGC CTGTCCCCAG ATGTCCCAGT	1200
10	AGCGATGAGT AACAGAGGTG GCTGTGGACT TCCTCTACTT CTCCTTGCTG GATCAGGGCC	1260
	TICCTGCCTC CCGCTGGGCA GGTCTGGCCT TGCTCTCTTG GCAGGGCCCC AGCCCCTCTG	1320
15	ACCACTCTGC AGCTCACCAT GCAGCTGATG CCAAAGTTGT GGTGTCCAGT GTGCAGCAGC	1380
	CCTGGGAGCC ACTGCCACCT TCAGAGGGGT TCCTTGCTGA GACCCACATT GCTTCACCTG	1440
20	GCCCCACCAT GGCTGCTTGC CTGGCCCAAC CTAGCGTTCT GTGCCATGCT AGAGCTTGAG	1500
20	CTGTTGCTCT TCTTCAGGGG AGGAAATAGG GTGGAGAGCG GGAAGGGTCT TGCTCCTAAG	1560
	TGTTGCTGCT GTGGCTTTTT TGCCTTCTCC AAAGACGCAC TGCCAGGTCC CAAGCTTCAG	1620
25	ACTGCTGTGC TTAGTAAGCA AGTGAGAAGC CTGGGGTTTG GAGCCCACCT ACTCTCTGGC	1680
	AGCATCAGCA TCCTACTCCT GGCAACATCA GGCCAACGTC CACCCCAGCC TCACATTGCC	1740
30	AGATGTTGGC AGAAGGGCTA ATATTGACCG TCTTGACTGG CTGGAGCCTT CAAAGCCACT	180
50	GGGATGTCCT CCAGGCACCT GGGTCCCATG ACCAGCTCCC CGTCTCCATA GGGGTAGGCA	186
	TTTCACTGGT TTATGAAGCT CGAGTTTCAT TAAATATGTT AAGAATCAAA GCTGTCTTTG	192
35	TTCAGGCTGC TATAACAAAA ATATAATAGC CTGGGTGGCT TAAAC	196

40 (2) INFORMATION FOR SEQ ID NO: 117:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 503 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 117:

50	AGTGATCCCC TTGCCTCGGC CTCCCAAAAT GCTGGAATTG TAAGCGTGGG CCTCTGCACC	60
	CGCCCTGGTC CGCAATTTAA AAACGCACAG CCACCATTCC CTYTCCAGAA AGCACCCAGA	120
55	TGCCTTTGGG AGAACCAGCC TCCTCCATGG AGGAAAGCTT GGGATCTGCC TTCCCACCTG	180
	GGGAGGAGAG GGATCTGTGG AAAATCCTTC TGACGGACTT CCCCTCAGTG CCTGATCCAT	240
	ACTCAATAGT AGAAAAAGTA AGAAATATAC AAAGATAGCA GATACACGGA GACAGTTCCC	300
60	CAAATAGCTG AGCGAWTAGC GCAGAAGCAA TATTGAAGAC CTAATAGCTG AGACATTTCC	360

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	AGAACTGATA	AAGTGCATCC	AGCCACAGAT	CAAGCAGCCC	AGAAAATTCC	AGGCAGCATC	420
5	AACAAATAAA	TAGCCCCACA	TGCACCCGTG	AAAATGCAGA	AGACCAAACA	AAAAAGTCCG	480
,	GTCAACAGCC	AGAGTTAAAG	AGG				503

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(2) INFORMATION FOR SEQ ID NO: 118:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1133 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 118:

60 GCCACAGCTT GGAATGAACC CCTGTGGATA AGGGGGACTA TTAGATAGAA TAAACATCAA TARATGCTTG ATGAATAAAC GCTAATCCTA CCTTCCCAGC CTGACACCTC CCAGTGGACA 120 25 CCACACTTCA CTTGAAGCCT TAGAAACCTT TCCCACCCAT GCTTCCAGCC CTGGCTTCAT 180 GTTGCCATTT CTCACCCCA GAACAGGCCG CCCGCCTGAA GAAACTACAA GAGCAAGAGA 240 AACAACAGAA AGTGGAGTTT CGTAAAAGGA TGGAGAAGGA GGTGTCAGAT TTCATTCAAG 300 30 ACAGTGGGCA GATCAAGAAA AAGTTTCAGC CAATGAACAA GATCGAGAGG AGCATACTAC 360 ATGATGTGGT GGAAGTGGCT GGCCTGACAT CCTTCTCCTT TGGGGAAGAT GATGACTGTC 420 35 GCTATGTCAT GATCTTCAAA AAGGAGTTTG CACCCTCAGA TGAAGAGCTA GACTCTTACC 480 GTCGTGGAGA GGAATGGGAC CCCCAGAAGG CTGAGGAGAA GCGGAACNTG AAGGAGCTGG 540 CCCAGAGGCA ANGAGGAGGA GGCAGCCCAG CAGGGGCCTG TGGTGGTGAG CCCTGCCAGC 600 40 GACTACAAGG ACAAGTACAG CCACCTCATC GGCAAGGGAG CAGCCAAAGA CGCAGCCCAC 660 ATGCTACAGG CCAATAAGAC CTACGGCTGT KTGCCCGTGG CCAATAAGAG GGACACACGC 720 45 780 TCCATTGAAG AGGCTATGAA TGAGATCAGA GCCAAGAAGC GTCTGCGGCA GAGTGGGGAA 840 GGGGCAGGG AGAGACAAGG CTGCTGCTAT TAGAGCCCCAT CCTGGAGCCC CACCTCTGAA 900 50 CCACCTCCTA CCACCTGTCC CTCAGGCTGG GGGAAAACAG GTGTTTGATT TGTCACCGTT 960 CGACCTTCGA TATCTCCCTC CCATCTCTCT CTCTCTCTGA GACTCTGAAT CCACAGCTCC 1020 55 GTATTTAATC TGTATTATTC CCCGTTCTTG GAATTTTCTT CCCATGGGGC TGGGGTACTT 1080 TACATTCAAT AAATACTGTT TAACCCAAAA AAAAAAAAA AAAAGAAAGA AGN 1133

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1	2	INFORMATION	FOR	SEO	ID	NO:	119:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1101 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 119:

	GGGCACAGCT	GAAGCTGCAG	ACCTCCCCAG	GGGATGGCTC	CTCTCCCCCA	GGAGCCCCGA	60
15	GGCAGGGGAG	GCAGAAAGCC	TGGGCTCTGG	GGGGTGGCCT	GCGGACAGCT	GIGCIGIGGG	120
	CCGGGGGCTG	GCCTGTCCC	ACAGGGNCGT	GGAGCTCGTG	GTTCTGAGCA	GCCAGCTGGG	180
	TGGTGTCTGG	GGATAGCTGG	GAGGCACAGC	GGCTGCCATG	TGGGACTGGG	ACTGGAGTGC	240
20	TCCCTGGTCT	TGGCCTCTGT	GGCTCAGCCT	TGCTCTGGTC	TGCCTGAGTG	CAGGGGCCAA	300
	GGGCACAGG	GCCAGTGAGG	CCGGCCACGC	TOGGGCCCTC	ACCTGTGAGA	TOGGGTCGGA	360
25	ATTTKACACA	GCCTANGGCT	TGGTTCTTGG	TKGTNGAMCG	TGGACTYCTK	AGAACGGGAG	420
	TGCTGGTCCT	.GAAAGGCGTG	GTTGGAGACC	AGCTGCTTTT	CTCGCTGTTT	TTCTCTTAGG	480
	AGATTAAACA	AAAACAGAAA	GCACAAGACG	AACTCAGTAG	CAGACCCCAG	ACTCTCCCCT	540
30	TGCCAGACGT	GGTTCCAGAC	GGGGAGACGC	ACCTCGTCCA	GAACGGGATT	CAGCTGCTCA	600
	ACGGGCATGC	GCCGGGGGCC	GTCCCAAACC	TCGCAGGGCT	CCAGCAGGCC	AACCGGCACC	660
35	ACGGACTCCT	GGGTGGCGCC	CTGGCGAACT	TGTTTGTGAT	ACTTOCCTTT	GCAGCCTTTG	720
	CTTACACGGT	CAAGTACGTG	CTGAGGAGCA	TCGCGCAGGA	GTGAGGCCCA	GGCGCCGAGA	780
	CCCAAGGCGC	CACTGAGGGC	ACCGCGCACC	AGAGCGTGAC	CTCGGCAGGC	TGGACACACT	840
10	GCCCAGCACA	GGCAGACCCA	CCAGGCTCCT	AGGTTTAGCT	TTTAAAAACC	TGAAAGGGGA	900
	AGCAAAAACC	AAAATGTGTG	ACTGGGCTTT	GGAGGAGACT	GGAGCCTCAG	CCCTGTCCTG	960
15	GCCACGGGCC	CCTCCCCCTC	CTCTCCCTCC	GCCTTGTGTG	CTGGATTTGT	AGCTTATCTT	1020
•	CCGTGTTGTC	TTTGGACCTG	TTTTAGTAAA	CCCGTTTTTC	ATTITAAAAA	AAAAAAAA	1080
	AAACTTTGGG	GGGGGGCCCC	N				1101

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(2) INFORMATION FOR SEQ ID NO: 120:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 282 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120:	
	AGCTTCTCTG TCCAGTCTTG AACTCTGGGS TCTCTTGGAA CTTTCCTCAC CCCTCTCAGC	60
5	CTGAATATTC CTTCCATGGA TTCCACTCAA CCAGACTTTG GATCTGTGCC TACTTAATCA	120
	ACCITATETT TGCAATATGT TCGGGCCCAC CTTCCACTCC TTGGTTCTTG TTCCTCCTTG	180
10	GCCTAACTTG TCCCTTCTCC ACTTCACATC CCCGGTGGGA CAGCATTCCT CCTTCCTCCC	240
10	AACCTCCCTC CGTCTCARAA AAAAAAAAAA AAAAAAAAA TT	282
15	(2) DECEMBER OF CO. TO VO. 121	
	(2) INFORMATION FOR SEQ ID NO: 121:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2635 base pairs	
20	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121:	
23	TAAGGGGGTG TGTGCTCACC TCCTCCTGAC CCTTAACACT CCTGTCCTGC CCAGACCAAC	60
	AGAGAGAGCT GTCCCTGAGA CCCCGGAGAG AAGCAGCTGC CGAAAGCTGC AGCCTTTCCG	120
30	CACTCTGAGA CCATGATCTT CCTCCTGCCA GGGGAGAGCC ACCCACAGGC CATGTCCAGC	180
	CCCACTTCCC TCAGCCCCCA GGGYTTCCTT CTGGCCCCTC TGAGGATTCC CTAGGGCTGC	240
35	CCCGCAGAGG GGYTTCCCCA AGCTCTGTTT TGAAGCCTGC AATGTGGAAA AGTGAGAAGT	300
	CAGAGGGAAC AGGACAGGTG CAGCCGGGCT CTGAGGCCAC ACCTCACACC TCGCTGTTCC	360
	CCAACATCCC CTGAGCAGTG TGAGCTCATC TCACCAGATG AGAAGAGGCC CTGTGCATTT	420
40	YTTTTGTTG TTTGTTGCTG TTTTCCCCCA CCCATCCAGT TCTCCTCAGC AAAGCAAATT	480
	CCTTAACACC TTTGGTGGAG AATTTCTTAC CCAGACTTGG GGCTGTGATG CCCTTCAGTG	540
45	CGTGGTGAGT GCAGCGTGTG TGCGTGTGCC TGTGTGTGAA CCTGGGGGCC ATCCTGGTGG	600
	CCTGGGAGCG TGAGGAGAGG CCCCCTGTGT GCTGGGTGAG TGGTGGGTGT GGGGTCAATG	660
-	CAGTGAGGCT CTCTGGGTGA GOCTCCCAAC CTGGCAGTCC CCAGCCTCCC AGCATCTGTG	720
50	AGCGTCTGTT GGACTTTACA GAAGAGCCTC ATCCYGTCTG CCCCTCACTC TGCCCTGGAA	780
	TCAACATCTT COGAGTCCTT CTTGGGGGAA ATAGCAGAGC CCCACTTAAC TCCATAAACT	840
55	SCTTCCCATT CCSCAGCCCA GITCTGATTG TTGAGGTGTC GCGTCGTTCC AGGTCCCCCA	900
	GTCCCCTCTT TCTCCTGTCC TCTCTCTGTC CTTCACCTCC CCACTCCAGC CCCGGCTCAG	960
	TTCAGGGAAA TOCTGTTCCA YATCAGCCCT CTGCTCTCTG AGGCAGCCGC GCCTCTGACT	1020

60 CGGAGCTACT TGAAACTTCT GCTCTTGCTA GGATTGGAGT CTACCTATCT CTTCCATTTG

	TCCCAGCTGG	AGTTCTGGAA	CTTTCCTCCT	CGGGGTGGGG	CICCCCCITTS	TTAAGGATGC	1140
5	TGGGGGGCCT	GGGGAAGGAA	GGAGTTCAGA	SGARGGGTGT	cccatattat	CTTSATGTCA	1200
	CCCTCCCCTC	CTGGGACACG	TECTCTCTCT	GICTCTGGGT	CTTCTGGGTG	TGCACGTTTG	1260
	TGTGTCCTTG	TAAATATGTT	TTAGGAAGAA	AGCAAAAGGG	ACTGARCTAG	CCTCTGGTAG	1320
10	GATTGCAGGG	GTCCAGCCTT	CCTCTTTCC	GAAGCCCCCA	CACTGCTTTT	CCCCCCACTG	1380
	AGACTGGTCC	CCTCAAAAGG	TAGACAAAAC	AGCAGCTCCC	TGTGGAGUTG	AAGGCCCCC	1440
15	TCAAAGTGGC	TTTTTGTTAG	ACAAGGITAA	GGTTTCCTCA	TGAGCAAGGT	TGCAGATCGG	1500
	TCCTTCCTCA	GCTCCTTGAT	TTGTGACCTT	GACCAAGGGG	CCTGCCACCC	AGCCCCTCCA	1560
	GTGCCCTCTC	CTCGATGCCT	CGCTCCTTCC	TGCCCCCACT	CCCCTGGTT	AGGEAGGTAG	1620
20	GGGAATTAGG	GCCATGCTGG	AAGAAGCTTA	ACCATGIGTT	CAAAGAACGG	TYTCYTGCTY	1680
	GCTTGGTCCT	GGAACTCCCC	TTGGCTGCCC	CAGGCCTCCT	TGGCCCATGG	GIGGIGGGG	1740
25	AGGTGGATGT	CAGATCTGGT	AGGTTGCAGC	AGAGAAAATA	AATGTGCCTT	GAGAGACCAC	1800
	TCAGAGAGGG	TCCAAGGGTG	ATGGAGAAGG	AAGCATGGCC	TOGGAGCTTG	GALGGGARGG	1860
	GIGGIGGGIG	GCGGCATCTT	GACTGCCCCC	TGTTGTCCCA	CACGTGGGGG	GTGGTCACCC	1920
30	CYCTTCACTC	CAGCCCGCCT	GCCTTCAGCC	TTCCATGAGC	TTCACCTGCT	TCCAACTTCA	1980
	CTTTGGAGGG	GGTGGGGTCC	GTTGGCATCA	ACACGGGGAC	CCTCTGCTTC	ACCAAAGCCC	2040
35	GAGCCCTCAG	CCCCTGGGGA	GAACAAATGG	CIGASCITIG	ATACCTGGGG	TCGTCGAGAG	2100
	GCTGCGGGCT	GGCGGCAGTC	CCAGGGGAGA	GACACCACAG	AAGGAGACCC	AGADATOCCG	2160
	AGGAAGTTCC	CAGCAGAGCA	AACTGCTTTC	CAGCCTGAAG	CCTGCTTAAA	CTGTGTGATG	2220
40	TGCAATAACT	GAGCTTAGAG	TTAGGAATTG	TGTTCALGTG	CTTGGATTTC	CGTCTGTAGA	2280
	TTTAACTGCT	GAAATTGTAT	CTCTCAGTAA	TTTTAGADGT	CTTTTAAAAA	ATTERARARC	2340
45	AAAGTGTTAG	ACTGTGTGCG	TGTGCGTTGA	TGGGCACTCA	AGAGTCCCTT	GASTCATCCA	2400
	GCCCTGCCTT	TCCCCTGCGC	CCCCATCCTC	TCACGTCCCG	ccciccatcc	ACTIGGGGAC	2460
	CCTGCCTCGT	GTCGTCTTTA	TCTGCCTATT	ACTCAGCCTA	AGGAAAC22G	TACACTOCAC	2520
50	ACATGCATAA	AGGAAATCAA	ATGTTATTTT	TAAGAAAATG	GAAAATAAAA	ACTITATAAA	2580
	CACCAAAAAA	далалалад	ACCCINGGGGG	GGGGCCGGTA	ACCCATTTCG	CCTAA	2635

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 994 base pairs

⁽²⁾ INFORMATION FOR SEQ ID NO: 122:

PCT/US98/11422

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WO 98/54963

(B)	TYPE: nucleio	acid
(C)	STRANDEDNESS:	double
(D)	TOPOLOGY: lin	ear .

5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 122:	
	GAATTCGGCA GAGGTTCGGC GAAGATAGGG AATAAGGAAG CACAGGAGTA GGGGAGAAGG	60
10	AAGCACAGGA GTAGGGGAGA TATACAGCGG TCAGGATAAG GGGGAAAGGG CGGTGGTTGC	120
10	SCAAGAGGTG AAACAAGATG TGAGAGACAA GGGGTAGGGA AGAAATGGGG CAGCGGTTAG	180
	GTTCAGAAGC GCATAGACCG TGGCGGACGG GCAATGCGAG GGGCACAGAA AGGAACTGAG	240
15	GGGTGGGCTA TTTTAARGGA GATGGTCCTT CAGCCCTCTT YTTTTCTGCG TAGTTCTCCT	300
	CCTCCAGGCC GCGCGCGGAT ATGTCGTCCG GAAACCAGCC CAGTCTAGGC TGGATGATGA	360
20	CCCACCTCCT TCTACGCTGC TCAAAGACTA CCAGAATGTC CCTGGAATTG AGAAGGTTGA	420
20	TGATGTCGTG AAAAGACTCT TGTCTTTGGA AATGGCCAAC AAGAAGGAGA TGCTAAAAAT	480
	CAAGCAAGAA CAGTTTATGA AGAAGATTGT TGCAAACCCA GAGGACACCA GATCCCTGGA	540
25	GGCTCGAATT ATTGCCTTGT CTGTCAAGAT CCGCAGTTAT GAAGAACACT TGGAGAAACA	600
	TCGAAAGGAC AAAGCCCACA AACGCTATCT GCTAATGAGC ATTGACCAGA GGAAAAAGAT	660
20	GCTCAAAAAC CTCCGTAACA CCAACTATGA TGTCTTTGAG AAGATATGCT GGGGGCTGGG	720
30	AATTGAGTAC ACCTTCCCCC CTCTGTATTA CCGAAGAGCC CACCGCCGAT TCGTGACCAA	780
	GAAGGCTCTG TGCATTCGGG TTTTCCAGGA GACTCAAAAG CTGAAGAAGC GAAGAAGAGC	840
35	CTTAAAGGCT GCAGCAGCAG CCCAAAAACA AGCAAAGCGG AGGAACCCAG ACAGCCCTGC	900
	CAAAGCCATA CCAAAGACAC TCAAAGACAG CCAATAAATT CTGTTCAATC ATTTAAAAAA	960
40	AAAAAAAAA AAAAAAAAAA AAAAAAGGGGA GGGG	994
45	(2) INFORMATION FOR SEQ ID NO: 123:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1542 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
50	(D) TOPOLOGY: linear .	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 123:	
55	GGCASAGCCA CCTCGGCCCC GGGCTCCGAA GCGGCTCGGG GGCGCCCTTT CGGTCAACAT	60
	CGTAGTCCAC CCCCTCCCCA TCCCCAGCCC CCGGGGATTC AGGCTCGCCA GCGCCCAGCC	120
	AGGGAGCCGG CCGGGAAGCG CGATGGGGGC CCCAGCCGCC TCGCTCCTGC TCCTGCTCCT	180
60	SCTSTTCSCC TGCTGCTGGG CGCCCGGCGG GGCCAACCTC TCCCAGGACG ACAGCCAGCC	240

	CTGGACATCT GATGAAACAG TOGTGCCTGG TGGCACCGTG GTGCTCAAGT GCCAAGTGAA	300
5	AGATCACGAG GACTCATCCC TOCAATGGTC TTAACCCTGC TCAGCAGACT CTCTACTTTG	360
J	GGGAGAAGAG AGCCCTTCGA GATAATCGAA TTCAGCTGGT TAMCTCTACG CCCCACGAGC	420
	TCAGCATCAG CATCAGCAAT GTGGCCCTGG CAGACGAGGG CGAGTACACC TGCTCAATCT	. 480
10	TCACTATGCC TGTGCGAACT GCCAAGTCCC TCGTCACTGT GCTAGGAATT CCACAGAAGC	540
	CCATCATCAC TGGTTATAAA TCTTCATTAC GGGAAAAAGA CACAGCCACC CTAAACTGTC	600
15	AGTOTTOTOG GAGCAAGCCT GCAGCCCGGC TCACCTGGAG AAAGGGTGAC CAAGAACTCC	660
	ACGGAGAACC AACCCGCATA CAGGAAGATC CCAATGGTAA AACCTTCACT GTCAGCAGCT	720
	CGGTGACATT CCAGGTTACC CGGGAGGATG ATGGGGCGAG CATCGTGTGC TCTGTGAACC	780
20	ATGAATCTCT AAAGGGAGCT GACAGATCCA CCTCTCAACG CATTGAAGTT TTATACACAC	840
	CAACTGCGAT GATTAGGCCA GACCCTCCCC ATCCTCGTGA GGGCCAGAAG CTGTTGCTAC	900
25	ACTGTGAGGG TCGCCGCAAT CCAGTCCCCC AGCAGTACCT ATGGGAGAAG GAGGGCAGTG	960
	TGCCACCCCT GAAGATGACC CAGGAGAGTG CCCTGATCTT CCCTTTCCTC AACAAGAGTG	1020
	ACAGTGGCAC CTACGGCTGC ACAGCCACCA GCAACATGGG CAGCTACAAG GCCTACTACA	1080
30	CCCTCAATGT TAATGACCCC AGTCCGGTGC CCTCCTCCTC CAGCACCTAC CACGCCATCA	1140
	TCGGTGGGAT CGTGGCTTTC ATTGTCTTCC TGCTGCTCAT CATGCTCATC TTCCTTGGCC	1200
35	ACTACTIGAT CCGGCACAAA GGAACCTACC TGACACATGA GGCAAAAGGC TCCGACGATG	1260
35	CTCCAGACGC GGACACGGCC ATCATCAATG CAGAAGGCGG GCAGTCAGGA GGGGACGACA	1320
	AGAAGGAATA TTTCATCTAG AGGCGCCTGC CCACTTCCTG CGCCCCCAG GGCCCTGTGG	1380
40	GGACTTGCTG GGGCCGTCAC CAACCGGGAC TTGTACAGAG CAACCGCAGG GGCCGSCCCT	1440
	CCCGNTTGTT CCCCAGCCCA CCCACCCCCT TGTTACAGAA TGTYTKGTTT GGGGTGCGGT	1500
45	TITIGIWATTG GITTINGGATN GGGGAAGGGA GGGANGGCGG GG	1542
	(2) INFORMATION FOR SEQ ID NO: 124:	
50	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1390 base pairs (B) TYPE: nucleic acid	
55	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 124:	

CAAGCTCTAA TACGACTCAC TATAGGGAAA GCTGGTACGC CTGCAGGTAC CGGTCCGGAA

	TTCCCGGGTC C	CACCCACGCG	TCCCCCCCTC	AGGGTGGACG	CATGGTTCTG	CACTGAGGCC	120
	CTCGTCATGG T	receccien	GTGGTACTTG	GTAGCGGCGG	CTCTGCTAGT	CGGCTTTATC	180
5	CTCTTCCTGA (CTCGCAGCCG	cccccccc	GCATCAGCCG	GCCAAGAGCC	ACTGCACAAT	240
	GAGGAGCTGG (CAGGAGCAGG	CCCGCTCGCC	CAGCCTGGGC	CCCTGGAGCC	TGAGGAGCCG	300
10	AGAGCTGGAG (GCAGGCCTCG	GCGCCGGAGG	GACCTGGGCA	GCCGCCTACA	GGCCCAGCGT	360
10	CGAGCCCAGC (GCCTGCCCTG	GGCAGAAGCA	GATGAGAACG	AGGAGGAAGC	TGTCATCCTA	420
•	GCCCAGGAGG A	AGGAAGGTGT	CGAGAAGCCA	GCGGAAAYTC	ACCTGTCGGG	GAAAATTGGA	480
15	GCTAAGAAAC 1	TGCGGAANINT	GGAGGAGAAA	CAAGCGCGAA	AGGCCCAGCK	TGAGGCAGAG	540
	GAGGCTGAAC (GTGARGWGCG	GAAACGACTC	GAGTCCCAGC	GCGAATGAGT	GGAAGAAGGA	6 00
20	GGAGGAGCGG (CTTCCCCTCC	AGGAGGAGCA	GAAGGAGGAG	GAGGAGAGGA	AGGCCCGCGA	660
20	GGAGCAGGCC (CAGCGGGAGC	ATGAGGAGTA	CCTGAAACTG	AAGGAGGCCT	TTGTGGTGGA	720
	GGAGGAAGGC	GTAGGAGAGA	CCATGACTGA	GGAACAGTCC	CAGAGCTTCC	TGACAGAGTT	780
25	CATCAACTAC	ATCAAGCAGT	CCAAGGTTGT	GCTCTTGGAA	GACCTGGCTT	CCCAGGTGGG	840
	CCTACGCACT	CAGGACACCA	TAAATCGCAT	CCAGGACCTG	CTGGCTGAGG	GGACTATAAC	900
30	AGGTGTGATT	GACGACCGGG	GCAAGTTCAT	CTACATAACC	CCAGAGGAAC	TGGCCGCCGT	960
50	GGCCAACTTC	ATCCGACAGC	CCCCCCCCT	GTCCATCGCC	GAGCTTGCCC	AAGCCAGCAA	1020
	CTCCCTCATC	GCCTGGGGCC	GGGAGTCCCC	TGCCCAAGCC	CCAGCCTGAC	CCCAGTCCTT	1080
35	CCCTCTTGGA	CTCAGAGTTG	GTGTGGCCTA	CCTGGCTATA	CATCTTCATC	CCTCCCCACC	1140
	ATCCTGGGGA	AGTGATGGTG	TOGQCAGGCA	GITATAGATT	AAAGGCCTGT	GAGTACTGCT	1200
40	GAGCTTGGTG	TOSCITOSTS	TOCCAGAAGG	CCTGGCCTAG	GATCCTAGAT	AAGCAGGTGA	1260
10	AATTTAGGCT	TCAGAATATA	TCCGAGAGGT	GGGGAGGGTC	CCTTGGAAGC	TGGTGAAGTC	1320
•	CIGTICTIAT	TATGAATCCA	TTCATTCAAG	AAAATAGCCT	GTTGCAAAAA	AAAAAAAA A	1386
45	AAAAACTCGA	•					1390

50 (2) INFORMATION FOR SEQ ID NO: 125:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1288 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 125:

60 GCCCCCCCC TGAAAGCCCC ATTGATCCAG CCTCCCCCCG CCTCCGAGCG CCGCCGASCA

	GACGCTGACC ACGTTCCTCT CCTCGGTCTC CTCCGCCTCC AGCTCCGCGC TGCCCGGCAG	120
_	CCGGGAGCCA TGCGACCCCA GGGCCCCGCC GCCTCCCCGC AGCGGCTCCG CGGCCTCCTG	180
5	CTGCTCCTGC TGCTGCAGCT GCCCGCGCCG TCGAGCGCCT CTGAGATCCC CAAGGGGAAG	240
	CAAAAGCCCC ATCCCGCAGA GGGAGGTGGT GGACCTGTAT AATGGAATGT GCTTACAAGG	300
10	GCCAGCAGGA GTGCCTGGTC GAGACGGGAG CCCTGGGGCC AATGGCATTC CGGGTACACC	360
	TOGGATCCCA GGTCOGGATG GATTCAAAGG AGAAAAGGGG GAATGTCTGA GGGAAAGCTT	420
	TGAGGAGTCC TGGACACCCA ACTACAAGCA GTGTTCATGG AGTTCATTGA ATTATGCCAT	480
15	AGATCTTGGG AAAATTGCGG AGTGTACATT TACAAAGATG CGTTCAAATA GTGCTCTAAG	540
	AGTTTTGTTC AGTGGCTCAC TTCGGCTAAA ATGCAGAAAT GCATGCTGTC AGCGTTGGTA	600
20	TTTCACATTC AATGGAGCTG AATGTTCAGG ACCTCTTCCC ATTGAAGCTA TAATTTATTT	660
	GGACCAAGGA AGCCCTGAAA TGAATTCAAC AATTAATATT CATCGCACTT CTTCTGTGGA	720
	AGGACTITIGT GAAGGAATTG GTGCTGGATT AGTGGATGTT GCTATCTGGG TTGGCACTTG	780
25	TICAGATIAC CCAAAAGGAG AIGCTICTAC IGGAIGGAAI ICAGIITÇIC GCAICATIAI	840
	TGAAGAACTA CCAAAATAAA TGCTTTAATT TTCATTTGCT ACCTCTTTTT TTATTATGCC	900
30	TTGGAATGGT TCACTTAAAT GACATTTTAA ATAAGTTTAT GTATACATCT GAATGAAAAG	960
	CAAAGCTAAA TATGITTACA GACCAAAGIG IGATITCACA IGIITITAAA ICTAGCATTA	1020
	TICATTITGC TICAATCAAA AGIGGITICA ATATTITITT TAGITGGITA GAATACTITC	1080
35	TTCATAGTCA CATTCTCTCA ACCTATAATT TGGGAATATT GTTGTGGTCT TTTGTTTTTT	1140
	CTCTTAGTAT AGCATTTTTA AAAAAATATA AAAGCTACCA ATCTTTGTAC AATTTGTAAA	1200
40	TGITAAGAAT TTTTTTTATA TCTGITAAAT AAAAATTATT TCCMACAACC TTAAAAAAAA	1260
	AAAAAAAA AAAAAAAA AAAAAAAA	1288

50

(2) INFORMATION FOR SEQ ID NO: 126:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1517 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 126:

AGTGGCTTAA AGGCATCGTT TTAGGGATTA CTGGGAAGTA TCTTCAAAGT AATACATGAG 60

AAACATTCCT TCCTAAATCC TTTATTATAT TGAATATCGT ATTAATTGGT TTTCAGAGGT 120

	TAAATTAACC ATGTATTCCT GCAATAAATG TCACTTGTMT CTTGTATATA ATCTTTTTTA	180
	TATATTACCG GATTGATTCA TTAGTATTTT GTTGAGGATT TTTGTGTCTA TATTCATAAG	240
5	AGATGCTGGT CTGCAGTTTT CTTTTTTTGT GATAATCTGG TTTTTGTATC AGTAATACAG	300
	GCCCCATGAA ACGAGTTGGG AAGTGTTCAC CTCTCTTGTA TTTTTTCAAG AGTTTGTGAA	360
10	GAATTGCTAT TAATTCTTTA AATGTTTGGT AGAATCTACC ATTGAAATCA TGTGTCCTGG	420
10	GCTTTTTTT GAGGGAAGTG TTCTGATAAC TAATTCAGTA TCTACTTTTT ATAGCTCTGT	480
	TCAGATTTIG CTTCTTCCTG AGITAGTTTT GGTAATTTGT GTATCTCTAG GARTTTGTCC	540
15	ATTICATITA TCTCATTIGT TGGCATAAAT TAAACTAAAT TTGGCCTGAG CCTACCTGTA	600
	TATCTTGAGT CCCTCTGTAA GGAACTGTAG CCTAACTTGT ACATAAACAA ACTGAAATCC	660
20	TAAATTAGGA ATGTAGTTTT TGTAACAGCT CCTGAGTCTC AGGCAGTCAC AGCAGYCAAG	720
20	TCTGTCAATT GCAGGCTGCT AACTAAGCAG CCCATGSTCA AATGAGGCAA AAACCTTTGC	780
	TTTTAACACA TAGTATAGCT TTGTAATCCT TTTCTTGCAC ACTCGGGTAA TTTCTTCCTT	840
25	TITCATICCC KGWATTITCC AKGAATATGA RICTYCCTIT TITCCCCTCC TGTCAGTCTA	900
	GCTAATGGTT TGTCAATTTT GTTGATCTTT TGAARAACAA ACCTTTGGTT CCACTTTCTT	960
30	GTTGCATATG CTGARTATTC TCATAATTGG AGTGGAAAGC TGATCTTTGA TTACTTATTT	1020
	TACTTAGGGC TGAGGAGTTC ATGGACTTCG CAAAACCTCC TTGAATCTAA ATTGCATCTT	1080
	CTTTCCTGGT TTCTGGGCTG AAACATGTTT TTTCCCATCT WANAWACCCT TGGTCTTTTC	1140
35	ATKGGCGATT AAGACTAGAG AAAGTTCTAG ATMCCTTGTC CTTTTATGCT GTCATTTTGT	1200
	TTAAAGGCTT TCTATGTAGT AAAACTATCT ATATAGACAA AATAGAGCCT TGAGITGTGG	1260
40	TCTTGAATTT GATCAACATG ATTTACCACA TTCTGTACTG GATATTTCTT CACCTGCTGC	1320
	TACTGTAAAC CATTTTATTC TTGGATCTTC TGTAGAGTAT ATTATCACAG GTACTTTTTA	1380
	CAGGGGTGTC TAATCTTTTG GCTTCCCTGG GCACATTGAA AGAAGAAGAA TTGTCTTGGG	1440
45	CCACACATCA AATACGCTAA CACTAATAAT AGTTGATGAG CTAAAAAAAA AAAAAAAAAG	1500
	GCAAAAAGN CCCAAAA	1517

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(2) INFORMATION FOR SEQ ID NO: 127:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1073 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 127:

	TGAATCTATT CTTTGAACAT TCTACAACAA GAATTACATT ATACTGTTAT ACCAGAGTAC	60
	TICTGCAGTG TGAAATAGAT TGGTTTGGAA AATGAACCTG GCTTTGCTAT AAATTACATT	120
5	CACAGGCCTT TTTGCAAATG TGTAACTTGC CTATCAAAGT AGTTTGTAGG GCAAATGCAG	180
	ANTATATGTC TCCATCTGGT ANAGTACCTT WINYTCATGT GGGANATCAN GTAGTATCAG	240
10	AACTTOGTCC AATAGTCCAA TITGITAAAG CCAAGGGCCA TICTCTTAGT GATGGGCTGG	300
	AGGAAGTCCA AAAAGCAGAA ATGAAAGCTT ACATGGAATT AGTCAACAAT ATGCTGTTGA	360
15	CTGCAGAGCT GTATCTTCAG TGGTGTGATG AAGCTACAGT AGGGRMGATC ACTCATGMTA	420
15	GGTATGGWIC TCCTTACCCT TGGCCTCTGW WICATATTTT GGCCTATCAA AAACAGTGGG	480
	AAGTCAAACG TAAGNTGAAA GCTATTGGAT GGGGAAAGAA GACTCTGGAC CAGGTCTTAG	540
20	AGGATGTAGA CCAGTGCTGT CAAGCTCTCT CTCAAAGACT GGGAACACAA CCGTATTTCT	600
	TCAATAAGCA GCCTACTGAA CTTGACGCAC TGGTATTTGG CCATCTATAC ACCATTCTTA	660
25	CCACACAATT GACAAATGAT GAACTTTCTG AGAAGGTGAA AAACTATAGC AACCTCCTTG	720
20	CTTTCTGTAG GAGAATTGAA CAGCACTATT TTGAAGATCG TGGTAAAGGC AGGCTGTCAT	780
	AGAGTTATGT GTTAGTCTCA GGAGTCTTAA CTTTTGAAAT ATGTTTTACT TGAATGTTAC	840
30	ATTAGATATT GGTGTCAGAA TTTTAAAACC AAATTACTGC TTTTTGAAAC CTCAAATTAT	900
	ATAATGTATC TTATGTATGT GCTTTATATT GTTATTTGTG TATACATTAA AATAATTCTG	960
35	AATTATTTAA TOTGATATGT TGTATTOTGT ATOTTGAAAT TTTTGTTTCC TTGAAACATG	1020
33	CATGCATTTA AAAATAAAGC TTAAACAACT GTAAAAAAAA AAAAAAAAAA	1073
40	(2) INFORMATION FOR SEQ ID NO: 128:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 300 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 128:	
50	CAACCCCTGC CTTTTTTTG TTTTCCATTT GCTTGGTAGA TCTTCCTCCA TCCCTTTATT	60
	TTGAGCCTAT GTGTGTCTCT GCCCGTGAGA TGAGTCTCCT GAATACAGCA CACTTACTGG	120
55	TCTTGACTCT GTATCCAATT TGCCAGTCTG TGTCTTTCAT TTGGAGCATT TAGCCCATTT	180
	ACATTTAAGG TKAATATTGT TATGTGTGAA TITRATCYTR TCATTATGWT GTTAGCTGGT	240
	TRANSPORTED CONSCIONATE CONSCIONAL ATOSTCTITA CARNITISCA	300

(2) INFORMATION FOR SEQ ID NO: 1	(2)	INFORMATION	FOR	SEO	ID	NO:	129) :
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1275 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

10

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 129:

GGCAGAGCCT GTCCCTGCTG CCCCTGCAAA AAAAACCCCC TCTGGTGTGA GCAGGATGGT 60 15 TOGAGGITAT GIGAGCICCT TCTCCTTTCC TCCAGTITCC TCTTCCCTTC TCCTCCCTGC 120 CTCTTTTGCT TTTCCCTTTC TTCCTGGTAC CCCCTGCCCA TTCCTGTATT TTCTCCCATC 180 GCCATTCTCC CCTCTCCCAC TGTCCCTAAC CCGTTCAAAC TCTTTCCTCT TAAATGGTTG 240 20 AGATTITICTO TOACCAAGOA CACCOCAGTA TTAATTAAAC TAGOTGCAAA CAGGCAGCAA 300 GTGGTCTACC ATGACAGATG GGTTTTGTGT GTGTGTGTGT GTGTGTAATT GTAATAAAAC 360 25 ATATTGARTC ACTCAATAAA CACAGAGTGT CTACTACATG TATCARGCAC TATCATAGAT 420 GCTAATTAAC GAAACTGAAA TGGCCAGGCC CTCACAGTGG CTCATGCCTA TAATCCCAGC 480 ACTITIGGAG GATGAGGCAG GAGGATCACT TGAGGCCGGG AGTTCAAGAC CAGCCTGGGC 30 AACATAGTAA GACTCCATCT CTACAAAAA AAAATTTTTT TTATTATACT TTAAGTTTTG 600 CGTTACATGT GCAGAACGTG TAGTTTTGTT ACATAGGTAT ATACGTGCCC TGGTAGTTTG 35 CTGCACCCAT CAACCCATCA CCTACATTAG GTATTTCTCC TAATGTTACC CCTCTCCTAG 720 CCCCCCACCC CGTGACAGGC CCTGGTGTGT GATGTTCCCC TCCCTGTGTC CATGTGTTCT 780 CATTGGTCAA CTCTCACCTA TGGAGTGAGA ACATGTGGTA TTTGGTTTTC TGATCTTGTG 840 40 ATAGCTTGCT GAGAATGTKG GTTTCCAGCT TTATCCACGT CCCTGCAAAG GGCATAAACT 900 CATCCCTTTT TATOGCTGCA TAGTGTTCCA TGGTGTATAC GTGCCACATT TTCTTAATCT 960 45 ATCATTGATG GACAAGTTTT GCTATTGTGA ATAGTGCCAC AATAAACATA CGTGTGCGTG 1020 TGTCTTTATA GCAGCATGAT TTATAATCCT TTGGGTATAT ACCCAGTAAT GGGATCACTG 1080 AGTCAAATGG TATTTCTCGT TCTAGATCCG TAAGGAATTG CCACACTGTC TTCCACAATG 1140 50 TTIGAACTAA TNIACACTCC CACCAACAGT GTAAAAGTGT TTCTATTTTT CCACAACCTC 1200 1260 TCCAACATCT GTTATTTCCT GACTTTTTAA TGAACGTCAT TCTAACTGGC GTGAGATGGT 55 1275 ATCTCATTGT GGTTT

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(2)	INFORMATION	FOR	SEQ	ID	NO:	130:
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5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 472 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 130:
	CNGAAACCCC GTGAACCCTC CCCGGGTTAA AAAGCCCCCC CTAAAT

CCCC CTAAATGGGG GGAACGCYTC 60 ACACGTTATA AAAAAGCACT AGAATGTTTT GAAAGCGAGA AACAACAGCT GTGTAGGGTA 120 GCTAGCAGTT AGTGTTGTAC AGAAGACAGA TATTTGTGCA TITYTGCATT TTCTAAGTTT 180 15 GCTGCAATGA GCATGTATTA CTTTCATAGT TATAAAACAC ATGCAAAATG CCCTTTTAAA 240 ATGAAAAAA ATCCATGAGT GTAAGTGATA TATATGCTTT GGAAAGCCTG GGACGGTCAT 300 20 TGTTTACTCT CAATAGTATG TGTTTGCCTT TGTCTTTTTG AGACATTTTG TTTTAATCTG 360 TTGATGACAA TAACCTGTTG ATAATATAAC TTGATAACAA ATAAAATGAC TTATGATTGA 420 ИИ АААААААА АААААААА АААААААА АААААААА ААА 472 25

30 (2) INFORMATION FOR SEQ ID NO: 131:

35

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1950 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 131:

40	ACCTCTCAGA ATCTTCTCTC AGCAACCTGA GTCTTCGCCG TTCCTCAGAG CGCCTCAGTG	60
	ACACCCCTGG ATCCTTCCAG TCACCTTCCC TGGAAATTCT GCTGTCCAGC TGCTCCCTGT	120
45	GCCGTGCCTG TNATTCGCTG GTGTATGATG AGGAAATCAT GGCTGGCTGG GCACCTGATG	180
45	ACTCTAACCT CAACACAACC TGCCCCTTCT GCGCCTGCCC CTTTNTGCCC CTGCTCAGTG	240
	TCCAGACCNT TGATTCCCGG CCCAGTGTCC CCAGCCCCAA ATCTGCTGGT GCCAGTGGCA	300
50	GCAAAGATGC TCCTGTCCCT GGTGGTCCTG GCCCTGTGCT CAGTGACCGA AGCTCTGCCT	360
	TGCTCTGGAT GAGCCCCAGC TCTGCAACGG GCACATGGGG GGAGCCTCCC GGCGGGTTGA	420
6 6	GAGTGGGGCA TGGGCATACC TGAGCCCCCT GGTGCTGCGT AAGGAGCTGG AGTCGCTGGT	480
55	AGAGAACGAG GGCAGTGAGG TGCTGGCGTT GCCTGAACTG CCCTCTGCCC ACCCCATCAT	540
	CTTCTGGAAC CTTTTGTGGT ATTTCCAACG GCTACGNCTG CCCAGTATTC TACCAGGCCT	600
60	GGTGCTGGCC TCCTGTGATG GGCCTTCGMA CTCCCAGGCC CCATCTCCTT GGCTAACCCC	660

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	TGATCCAGCC TCTGTTCAGG TACGGCTGCT GTGGGATGTA CTGACCCCTG ACCCCAATAG	720
_	CTGCCCACCT CTCTATGTGC TCTGGAGGGT CCACAGCCAG ATCCCCCAGC GGGTGGTATG	780
5	GCCAGGCCCT GTACCTGCAT CCCTTAGTTT GGCACTGTTG GAGTCAGTGC TGCGCCATGT	840
	TGGACTCAAT GAAGTGCACA AGGCTGTGGG GCTCCTGCTG GAAACTCTAG GGCCCCCACC	900
10	CACTOCCCTG CACCTGCAGA GGGGAATCTA CCGTGAGATA TTATTCCTGA CAATGGCTGC	960
	TCTGGGCAAG GACCACGTGG ACATAGTGGC CTTCGATAAG AAGTACAAGT CTGCCTTTAA	1020
15	CAAGCTGGCC AGCAGCATGG GCAAGGAGGA GCTGAGGCAC CGGCGGGCGC AGATGCCCAC	1080
13	TCCCAAGGCC ATTGACTGCC GAAAATGTTT TGGAGCACCT CCAGAATGCT AGAGACCTTA	1140
	AGCTTCCCTC TCCAGCCTAG GGTGGGGAAG TGAGGAAGAA GGGATTCTAG AGTTAAACTG	1200
20	CTTCCCTGTT GCCTTCATGG AGTTGGGAAC AGGCTGGGAA GGATGCCCAG TCAAAGGCTC	1260
	CAAGCGAGGA CAACAGGAAG AGGGATCCAC TGTTACCAAA AGTCCTGATT CCCCCATCAC	1320
25	CAACCTACCC AGTITGTTCG TGCTGATGTT GGGGGGAGATC TGGGGGGAGT TGGTACAGCT	1380
	CTGTTCTTCC CTTGTCCTAT ACCGGGAACT CCCCTCCAGG GTACCCACAG ATCTGCATTG	1440
	CCCTGGTCAT TTTAGAAGTT TTTGTTTTAA AAAACAACTG GAAAGATGCA GAGCTACTGA	1500
30	GCCTTTGCCC TGAATGGGAG GTAGGGATGT CATTCTCCAC CAATAATGGT CCCTCTTCCC	1560
	TGACGTTGCT GAAGGAGCCC AAGGCTCTCC ATGCCTTTCT ACCTAAGTGT TTGTATTTTA	1620
35	TTTTAAATTA TTTATTCTGG AGCCACAGCC CCCTTGCTTA TGAGGTTCTT ATGGAGAGTG	168
	AGAAAGGGAA GGGAAATAGG GCACCATGGT CCGGTGGTTT GTAGTTCCTT CAAAGTCAGG	174
	CACTGGGAGC TAGAGGAGTC TCAAGCTCCC CTTAGGAAGA ACTGGTGCCC CCTCCAGTCC	180
40	TAATTTTTCT TECCTECCCC GCCTTEGGGA ATGCCTCACC CACCCAGGTC CTGACCTGTG	186
	CAATAAGGAT TGTTCCCTGC GAAGTTTTGT TGGATGTAAA TATAGTAAAA GCTGCTTCTG	192
45	TCTTTTTCAA AANAAAAAA AAAAAAAACT	195

(2) INFORMATION FOR SEQ ID NO: 132:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 990 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 132:

TOGAAGATIT AAAATAGGIT TCATATITCT CTTGAATATG AATATATAAG CTTGAATAAG

50

	CHECAGECCI TACTATIATE ARATTITCCT TATTATITCT ACCAATGCTT CITATATAA	120
	ACCORDATES THUTCHIATE ASSAURATED CATTAGORGO CROTOGATEA ACATTROCAT	180
5	SAARGIATT TYTECATTET TISATCITAA ACTITITGIG TCTTTATATA AGGIATGCTY	240
	CTECTAASCA TGATATITTT AACCACAATA GITGAAAGAC AATCTYCACC TTTTACTTGT	300
	ATAITTACAI GTAAFFIAAF TIITGATGCA TATTACGTCT TATTATTTAA CCAACCTATT	360
10	TTATTITATO TAGGGCATTT TYCAGAAAGC CITATTITCT TGTATTAATC AAATATTTT	420
	AYCAPIGTAT TYTOCYCTAT TASTTAGKAA TACGKTACYC YAAATATATA TYGYGGSTAT	480
15	TTTCAGAATT GCAATATGCC TCCTTAATTT ATTAGAGGCT AACCTAAATT ATTACTTTTA	540
	CCACTTACTT GAAAATTCTG GAACTTTAGA ACATTTATTG TTTTATGCAT TTTAATTCTA	600
	CTTGTATITT TACTACTOCT ALACATTATT ATTGTTTTAG ACAAGCCAAA ATATATNITG	660
20	TTACTATOTT ATYCICCATT TOTTTCTGTA TTTTTATGCC ACTATGTATG CTCAATTTCC	720
	TTCTATGTGA TGAACCTAAT TCAGTACTTT TGTTTTTTAA TCTGTGCAGG TAGCCTGGCC	780
25	ATTAAATITI TATTIYIGGI TIGCIGAAAA AATTGIGITI ATTICIATAT GCATACITAT	840
	SCAPATAGAA TACTAGGING ACATATTITT AGTATTTATA AATGTAAAGT CATTWATTKG	900
	GCTICTATCA THICKGIKGA GAAATCAATT GTCAGCCCAA TAGTTTTTCA TTTTAAATTA	960
30	CNGAATTITT TCATGTOTOT GGTTTTAGGA	990
35		
	(2) INFORMATION FOR SEQ ID NO: 133:	
	(i) SEQUENCE CHAPACTERISTICS: (A) LENGTH: 1720 base pairs	
40	(3) TYPE: nucleic acid (C) STRANCEDNESS: double	
	(D) TOPOLOGY: linear	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 133:	
45	GTCTGACAAG CGACTGTGGT TATTCCCCTA AAGTTTACTT CAGCACTAAC ACTAGTGCTT	60
	CCGCTGGAGT TTGCAGTTTT CCAGCTTTAT ACAGGATTTT CCTTTGACTG GAAGAGTCAA	120
50	GGATATAGAG ACTCAACAGT GACATITATT GTACAACATC AAGGGGAATA GGATACTCAT	180
	CARACTOGGA TTATTCTTAT CHARACATOG TCTTCTTTGA ATANGAAAAA TACATAGTTG	240
	GITATTATGG ACTTALAACT GIGTTAAATG GATATTCTGA TAAAATATTT GCTGCTCTGT	300
55	AGASTGTGGA AAATCTSAGA ASATTAGCTT TACTCATCTT GAGCTTTGAG GATGTTCTCT	360
	GIRCGCCGAT GGITTCHTAT TARCTARARA AGCTGGGTAT TGTARARTCT CATTTATARA	420
60	AACTCAGATG AGAAGAAAAT TITCTTTGAT GGTGAGACTG TTGTCTTAGT TCAGGAAATT	480

	ATTTAATAAT CCTTTGTTAC CTGTGAATGA AGGAACTTTG TAATTCTGAT TTATCGTAAA	540
_	ACATGAGCCT TTCCAGAGTC AGCTTAGACA CTGTTGTCGC AAATAGCCAT GCTTTGCCTT	600
5	ATGCCAAGGA GGCCCAGAGG GAGGGCCTAG TCTTCCTCTG TTGCTGTACA TATATTGAAA	660
	TECTTTTTT TTTTATTTTG CATTTGTTAT CTATAATGAG CTTTCTGAGC CCTGATATTA	720
10	TGTGAGACAA ACAGGAGTTA TTGATGTTAT ACACTCCCTT CCATTCAGGA TTTTCTGCTT	780
	GGAGGGAAAT ATGTTGACCT TAGAGAATTG TGAATATTGT TGCAATTCTT GAATATATTA	840
	CCATGTGAAT AATAGAGACT GTGTTGCTCT CTAGTATAAG CTATATTTAT TTTTGATTCA	900
15	TTTGAATTAC TAGTTATAAC TGGAGAAATT TTGTTACCTC TATCCTGGCT TGCCTGACTG	960
	GCTGTATAAT AGCAGCAGCC TCTTTTAGAG CATCTTAATG AAAACATGGA TGAAAGGAAT	1020
20	TAATGATGAT ATCTGCAGAC TGCGTAGAAA ATGGCTTTTG TTCCCAGCGT TAACATTTTC	1080
	TTCTCAATCA CATTTCAATG TTTGTGGAGA GTGGCAGATT CACACCAGAA ACACTAGGTG	1140
25	TTCATATCCA TAGCATGGAT GCAGAATAAG CAGTTGGGAG AGAAGCTTCT TCCTACCTGG	1200
25	TACTCCTCCC ATTCACCTCA GCCCAGCCCC AGACAGGCGT TAGCATTCAG TGTGGGCCCT	1260
	CAGGCAGCCC TGAAGCCTGG CTGGGTCATC AGATGGGGGC AGCCTGTGAC GGGCACCAGC	1320
30]	GCCCTGATTC CAGGGAAGAG TTCCTGGAGG GTGTTGGCTG TTTTTGTTAG CTCAGTTTTT	1380
	TTCTGGGCTC CACCATTCCT AACTCCAGGT AGACAAGATA GATGTCACAC ACAACAATTT	1440
25	TAAAGTATTT TGCTTAGTGC ATTTTGTTTA TGATTGCAGT GTTTGTTTCT TATTTAATAG	1500
35	GCTTTTTACT TCATTCTATT AAATTTTAGT GTTTAGAAGA GGCGGGTACT GTCACTGTGT	1560
	AAAATATGTA ATATTTTATA TGTTATACCA TGTCATATAT ACTTGCAATA TCAGACCTTG	1620
40	CATTCAATAT ACAATGCAAT TGACTCTTTG CAGACCTGCA TTTTTCAGTG AACAATAAAA	1680
	AGATTGTCTG GCACTCCAAA AAAAAAAAA AAAAAAAAAA	1720

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(2) INFORMATION FOR SEQ ID NO: 134:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 705 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 134:

GGCACGAGGC CATCTGGGCT CATTCAGCAG GAAATAATGG AAAAAGCTGC AATATCCAGG 60
TGTTTACTAC AATCTGGAGG CAAGATCTTT CCTCAGTATG TGCTGATGTT TGGGTTGCTT 120

	GTGGAATCAC AGACACTCCT AGAGGAGAAT GCTGTTCAAG GAACAGAACG TACTCTTGGA	180
	TTAAATATAG CACCTTTTAT TAACCAGTTT CAGGTACCTA TACGTGTATT TTTGGACCTA	240
5	TCCTCATTGC CCTGTATACC TTTAAGCAAG CCAGTGGAAC TCTTAAGACT AGATTTAATG	300
	ACTCCGTATT TGAACACCTC TAACAGAGAA GTAAAGGTAT ACGTTTGTNA AATCTGGGAA	360
	GACTIGACIG CTATICCATI ITGGGTATCA TATGTACCIT GATGAAGANG ATTAGGTIGG	420
10	GATACTICAA GIGAAGCCTC CCACTGGAAA CAAGCTGCAG TIGITTTAGA TAATCCCATC	480
	CAGGITGAAA TGGGAGAGGA ACTIGTACTC AGCATTCAGC ATCACAAAAG CAATGTCAGC	540
15	ATCACAGTAA AGCAATGAAG AGCAGTTTTC CAATGAAAAC TGTGTAAATA GAGCATCAAC	600
	AAGTACAAAA TTCTTGTCTT AATTAGTGGG GGTATATAAA AATTCCTTGT AATGGTCAAA	660
	TATTITITAA AATTGACATT AATAAAGCAT ATTTTAAAAG TTTCT	705
20		
25	(2) INFORMATION FOR SEQ ID NO: 135:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 323 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 135:	
35	AGCACACAC TCCTTTAGTT GCTCCTAAGG TCATGTTCAA CATTCGTGGA GTGCATTTTC	60
23	TGCTCAGGGA GCTTTCCCAG ACCCGGAATG TTTGGTGCTC ACAGACYCTG GCAAGGATCG	120
	GTATTGCTGT TCCTCAGTTT TGCCTGGGGA AATGGAGGST CAGTGACGTT CAGTGACGTG	180
40	CCCAGAGTCA TGCCATTGGC GGGTGGCCCA GKGMTCCAGG TCTCCAGCAC CCCTCGGCCC	240
-	CCTCCTCACC AGGTCACATC ATCTCCTGGA TTAGAATCTG CTCACATAGT CTGTCCTGAA .	300
45	AGGAAAAAA AAAAAAAAA AAC	323
43		
50	(2) INFORMATION FOR SEQ ID NO: 136:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 582 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 136:	
	GGACGGAATG GTGCAACCCT CCTWAMTTTT CTKGKGCTGT TGACAACAGA GGGACGGACG	6

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	GAAAACATTT TTYGTGGGAG AATCCTACYT CTGCAGSGGA GCCCTTAAGC GATKGATTTT	120
	GAATCTKGAC CCTTTACCAA CTAATTTTGA AGGAAGATAC CTTGGAAATA TTTGGCATTC	180
5	AGTOGGTTAC TGAAACAGCA TTAGTGAATT CATCTAGAGA ACTCTTTCAT TTATTCAGGC	240
	AACAACTGTA CAACTTGGAA ACCTTGTTAC AGTCCAGTTG TGATTTTGGG AARGTATCAA	300
١٨	CTCTACACTG CAAAGCAGAC AATATTAGGC AGCAGTGTGT ACTATTTCTC CATTATGTTA	360
10	AAGTTTTCAT CTTCAGGTAT CTGAAAGTAC AGAATGCTGA GAGTCATGTT CCTGTCCATC	420
	CTTATGAGGC TTTGGAGGCT CAGCTTCCCT CAGTGTTGAT TGATGAGCTT CATGGATTAC	480
15	TCTTGTATAT TGGACACCTA TCTGAACTTC CCAGTGTTAA TATAGGAGCA TTTGTAAATC	540
	AAAACCAGAT TAAGGTTTGA CTGGTTTCAT TTGATTTTTA AG	582
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20	(A) TIMOPHUMYOU POP GTO XP NO. 127.	
	(2) INFORMATION FOR SEQ ID NO: 137:	
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1021 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 137:	
	TTCGGCAGAG CCCTTGCGCG CTCTTGAATA CCTGCKTTCT GTAGCGCTAG TTCTCTTCAA	60
	GATTICCTTA GIGTCATTIC ATTICGGTTT CTTTTCTCGC CATGTTTTC TGTCGGAATT	120
35	ACGGTTCGTT TTGGTTCTAT GTACTCTCTA AAATGTTATC GTTTTTCATT TGTCTACTAA	180
	TTTTCGTGCA TTTGTTACTA CTGAGTTTCT TAATATCTGA CTGGCCTCCG CCCACGGCT	240
40	CTGCAGANCA TAAAATACTC AGGCTGATGG TAGTGCAGAG ACTCTCCCTC CTTGATCAGC	300
	GCAAACGTTG GTCTGAGGCT TGAGGGATGG AGCAACATTT TCTTGGCTGT GTGAAGCGGG	360
45	CTTGGGATTC CGCAGAGGTG GCGCCAGAGC CCCAGCCTCC ACCTATTGTG AGTTCAGAAG	420
43	ATCGTGGGCC GTGGCCTCTT CCTTTGTATC CAGTACTAGG AGAGTACTCA CTGGACAGCT	480
	GTGATTTGGG ACTGCTTTCC AGCCCTTGCT GGCGGCTGCC CGGAGTCTAC TGGCAAAACG	540
50	GACTOTOTOC TOGAGTOCAG AGCACOTTOG AACCAAGTAC AGCGAAGCCC ACTGAGTTCA	600
	GTTGGCCGGG GACACAGAAG CAGCAAGARG CACCCGTAGA AKARGTGGGG CAGGCAGARG	660
.	AACCOGACAG ACTCAGGCTC CRGCAGCTTC CCTGGAGCAG TCCTCTCCAT CCYTGGGACA	720
55		780
	GACAGCAGGA CACCGAGGTC TGTGACAGCG GGTGCCTTTT GGAACGCCGC CATCCTCCTG	760

60 TIGGITTIGC CCCGITCTCT GTACTCTCGG CGTCCTGTTC ACGGATCTGT GGAGCTAAGC

AGCCTTAGAT AGCAGCAGAA GGCTTTTTGG ATTCTCCTCC TTGAAAAGAT TCTCAGTTAC 960

CAAACGTCTC CACCTAGAAA ATAAAAATAC ATTAAGATGT TGANAAAAAA AAANAAAAAA 1020

A 1021

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(2) INFORMATION FOR SEQ ID NO: 138:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1777 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 138:

CGGAAGATGA TGGCTTCAAC AGATCCATTC ATGAAGTGAT ACTAAAAAAT ATTACTTGGT 60 ATTCAGAACG AGTTTTAACT GAAATCTCCT TGGGGAGTCT CCTGATCCTG GTGGTAATAA 120 25 180 GAACCATTCA ATACAACATG ACTAGGACAC GAGACAAGTA CCTTCACACA AATTGTTTGG CAGCTTTAGC AAATATGTCG GCACAGTTTC GTTCTCTCCA TCAGTATGCT GCCCAGAGGA 240 TCATCAGTIT ATTITCTTTG CTGTCTAAAA AACACAACAA AGTTCTGGAA CAAGCCACAC 300 30 AGTCCTTGAG AGGTTCGCTG AGTTCTAATG ATGTTCCTCT ACCAGATTAT GCACAAGACC 360 TAAATGTCAT TGAAGAAGTG ATTCGAATGA TGTTAGAGAT CATCAACTCC TGCCTGACAA 420 35 480 ATTCCCTTCA CCACAACCA AACTTGGTAT ACGCCCTGCT TTACAAACGC GATCTCTTTG AACAATITCG AACTCATCCT TCATTTCAGG ATATAATGCA AAATATTGAT CTGGTGATCT 540 CCTTCTTTAG CTCAAGGTTG CTGCAAGCTG GGAGCTGAGC TGTCAGTGGA ACGGGTCCTG 600 40 GAAATCATTA AGCAAGGCGT CGTTGCGCTG CCCAAAGACA GACTGAAGAA ATTTCCAGAA 660 TTGAAATTCA AATATGTGGA AGAGGAGCAG CCCGAGGAGT TTTTTATCCC CTATGTCTGG 45 TCTCTTGTCT ACAACTCAGC AGTCGGCCTG TACTGGAATC CACAGGACAT CCAGCTGTTC 780 ACCATGGATT CCGACTGAGG GCAGGATGCT CTCCCACCCG GACCCCTCCA GCCAAGCAGC 840 CCTTCAAGTT CTTTTATTTC TGGGTAACAG AAGTAGACAG ACAGGTTACT TGGTGTATCT 900 50 TCTGTTAAAG AGGATTGCAC GAGTGTGTTT TCCTCACACA CTTTGATTTG GAGAATTGGT 960 GCTAGTTGGC AATAGATAAC TCAGCGTAGA TAGTATTGCA AAAAGGGGAG GAAATACACA 1020 55 1080 ACAATAATAA ATGTAAAAAC CTGCTATTCA ACATGCAGTT TTATTTCGAR GCCAAAAATC TAGAGCTITIC CCAAGATCCT GITGCCTTAG GCACATNCAC ACTTCAACAG TGCACACTAT 1140 CCAACAGTGC ACACTATTCA ACAGTGCACA CTATTCAAAA GCGTAGACTA TTTTTTTGCA 1200 60

•	TGTTCAAGAT	ATTTGTTTTG	GTCTTATGTG	TGTGTGAGAG	AGAGAGATTC	CTTTGACATT	1260
	AAGGAGCATC	AATGAGAAAA	GATGATGAGG	CAGGAATTAA	TAAAGAAATG	AAGTCGTGTG	1320
5	TGTTTGGTTG	CCTGTCAGAG	GGCACACAAT	TTCATAAACA	CCATGCCTGG	ACAATTTGAT	1380
	ATTAATATTT	AACACCTCTG	CATCTTTTTC	TTAAAAAAGA	ATATGGGCCA	GATACAGTGG	1440
10	CTCACATTTG	TAATCCCAGC	ACTTTGGGGA	GCCAAGTTAG	CAGAATCCCT	TGAGCACAGG	· 1500
10	AATCTGAAAC	CAGCTTGGGC	AACATAGTGA	GATCCCATCT	NTACAAAAA	CTTAAAAATT	1560
	AGCCAGGCAT	GATGGCACAT	TCCTGTAGTC	CTAGCTACTC	AGGAGGCTAA	GCTAGGAGGA	1620
15	TTGCCTGAGC	CCAGGAGTTC	AAGGCTGCAG	TGAGCTAAGN	ACGTGCCAGT	ACACTCCAGC	1680
	CTGAGCCACA	AAGTGAGACC	CTGTCTCGCA	ИАААААААА	TTAAAAAGTC	GCGCGCGCC	1740
20	CCGGTACCCA	AATCGCCGGA	TATGATCGTA	AACAATC			1777

(2) INFORMATION FOR SEQ ID NO: 139:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 643 base pairs (B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139:

	TTTTTTTTT	TTTTTTTT	TTTTTTTTT	TTTTTTTGGG	aatgagaaaa	TAACTTTATT	60
	TTCATTGTGG	GGAGCGGGCC	GATGTCCAGC	CTCAGAACTT	CTGGAACTGC	TTCTTGGTGC	120
	CGGCAGCCTT	GGTGACCTTG	AGCACGTTGA	AGCGCACTGT	CTTGCTCAGA	GCCCGCCACT	180
	CCCCACTGT	GACGATGTCA	CCGATCTGGA	CCTCCCTGAA	GCAGGGGGAC	AGGTGTACAG	240
	ACATGITCIT	CICCCCITC	TCGAAGCGGT	TGTACTTGCG	GATGTAGTGC	AGATAGTCTC	300
	GGCGGATGAC	AATGGTCCTC	TGCATCTTCA	TCTTGGGTCA	CCACGCCAGA	GAGGATCCGC	360
	CCTCGAATGG	ACACATTACC	AGTGAAGGGG	CATTTCTTGT	CAATGTAGGT	GCCCCTCAAT	420
	ACCTCCTTG	GGGTGTCTTT	GAAGCCCAGA	CCGATGTTCT	TGTTAGTAAC	CCGCGGGAGC	480
١	TTCTCCTTGC	CAGTTTCTCC	CAGCAGGACC	CICTICTIGT	TTTGAAAGAT	GGTCGGCTGC	540
	TTTTGGTAGG	CACGCTCAGT	CTGAATGTCC	GCCATCTTCT	CGTGCCGMAY	TCCTGCAGCC	600
	CGGGGGATCC	ACTAGITCTA	GAGCGGCCGC	ACCGCGGTGG	AGC		643

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(i) SE	OUENCE	CHARACT	TERISTICS:
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(A) LENGTH: 1220 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 140:

10	GGCACGAGGA	TGATAGACCT	ACTGGAGGAA	TACATGGTTT	ACAGGAAGCA	TACCTACATR	. 60
10	AGGCTTGATG	GCTCATCCAA	GATCTCGGAG	AGGCGAGACA	TGGTTGCTGA	TTTTCAGAAC	120
	AGGAATGACA	TCTTTGTGTT	CCTGTTAAGC	ACACGAGCTG	GAGGACTGGG	TATCAATCTC	180
15	ACTGCTGMAG	ACACAGTGCA	TTTTCTATGA	TAGCGACTGG	AACCCCACTG	TGGACCAGCA	240
	GGCCATGGAC	AGGCCCACC	GCTTAGGGCA	GACAAAGCAG	GITACTGTGT	ACCGCTCAT	300
20	CTGTAAAGGC	ACCATTGAAG	AACGCATTCT	GCAAAGAGCC	AAGGAGAAGA	GTGAGATTCA	360
20	GCGGATGGTG	ATTTCAGGIG	GGAACTTCAA	ACCAGATACC	TTGAAACCCA	AAGAGGTGGT	420
	TAGTCTTCTT	CTAGACGACG	AAGAGTTGGA	GAAGAAACGT	ATGTACTCTA	AACCTCTATA	480
25	CACTCCCCTC	ACGTATCTGA	GAATGGAAGA	GGTACTTGGS	TGTGTGCCAA	GGGTTAGGCA	540
	AAGCCAGAGG	CTGTATTTAG	GGAAAGTATT	TTTGTGCTCA	TATTTTATAT	AAAAACCCAA	600
30	ACAAGAATGT	GTTTGTAGGC	CAGGCGTGGT	GGCTCGCGCC	TCTAGTCTCA	GCATTTCGGG	660
50	ARGCCAAAGT	GGGCAGATCA	CCTGARGTCA	GGARTTTGAG	TTTGARACCA	GCCTGGCCMA	720
	CGTTGTGAAA	CCCCACCTCT	ACTARGARTA	CSGAAAATTG	GTTGGGCATG	GTGGCGGGCA	780
35	CCTGTAATTC	CAGCACTITG	GGAGGCTGGG	GCAGAANAAT	TGCTTGAGCC	CAGGAGGTGG	840
	AGATTGCGGT	GAGCCGAGAT	YGTÇÇCATTC	CAMPCCAGCO	: SGGGCAATAA	GAGTGAAAYT	900
40	CCATCTTTTA	AAAACAAACA	AAAACAAAA	ACACAAGACG	GCTCACACCT	GTAATCCCAG	960
70	CACTITIGGGA	RGCCGARGCA	GCTGGATCAC	: GARGTCAGGA	GTTCCAAGAC	TAGCCTGGCC	1020
	AACCIGGIGA	AGCCCCGTCT	CTACTAAAA	TACMAATATI	AGTCGGGCGT	GTGGTGGC	1080
45	ACGTGTAATC	CCAGCTACTC	GGGAGGCTG	A GGCAGGAGAA	TCCCTTGAAC	CTAGGAGGCA	1140
	GAGGTTGCAG	TGAGCCAGG	1 TOGTGCCATT	CACTCCAGO	CTGGACAAC	AGAGCAAGAT	1200
50	TCCATCTCAA	KAAAAAAA	.				1220

(2) INFORMATION FOR SEQ ID NO: 141:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 721 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(XI) Sequence Description. Seq 15 no. 1-1.	
_	AATTCGGCAC GAGCCAGGTT AGCCGGAAGG GCAGCTCTCC AGGCCCTGCC CACCCCACAG	60
5	GGGGCTCCTT ATGCACAGCG GGGCGTCTCC TTGTGGCCAT AGAAACGGAA CTGGCTCTTT	120
	TCAACAGTGC TGCAAGAGGA TGGTTATTTA ACGCTGGCCC CCAAGGAGGA AAGGCACAGA	180
10 15	CYTTCCTCCC TCCTGGAACA TCCAAGGGCA CTGGATCCTC TGTGTCCCTC TGAGATGGGG	240
	TOCCACTCCA GCAAGAGCAC CACGGTGGCA GCTGAGTCCC AGAAGCTTGA AGAAGAGYGC	300
	GAGGGAAGAG AGCCAGGTCT GGAGACCGGC ACCCAGGCAG CAGACTGCAA GGATGCCCCG	360
13	CTGAAGGATG GAACCCCTGA GCCAAAGAGC TGAAATGCCT CTCTCCAGAG TCGGACCCTC	420
	ACCTCYTTCC TGGAACTGCC TTTGGCCCCA GAACCATGAG ACAATCCCCA CCCTGAGAAG	480
20	CTCCGATCAC TGGGAGGAGA GAGAAAGCCT CCAGCTTTGG GATTCAGGCT TCAGAAGTTT	540
	TTAGCAGCCT TTGCTCATTG GAGAGGTGGG GAAAGGATAA AGTTCTTATA AGGAAATCCC	600
25	TAATTTCCCC CAGCTCCTCC CCNCCNGAAG AAGGAACNAA AGAAAGTTCC TTCCACACGT	660
23	TITGTTGGAA ACTITTCCCT TGCCAACTIT CCTTGGATTG CCAGAACAAA GCCCTCCAGA	720
	A .	721
30		
	(2) INFORMATION FOR SEQ ID NO: 142:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1468 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:	
•	ATGAATTAAT GTTTATAAAT GACTGTACTG AATTTAAAAC CGTACAGTTT CATTTGCATT	60
45	TTGACATTAC TTTATTATAC ATTTTGCATT TAAAAGGCTG CACCAGTTGG CTTTTCTTCT	120
	GTTTTATTCT CAAAATATAG AGATTCTGTG ATTTATTTGC CCTGTTTATG GATTAAAAAG	180
50	AAAATTCTAA TATAAAGCAT TTCAATAGGA TGCATAGGTA TATTACGTTT TTTAAATGCT	240
50	TTAGATCTGT GATTCTTGAC TTACTATITA TTTTATCCCC TTTAAGTCAG GGATGCTTTA	300
	TICTATTITA AAGCACTTAT GAGITACATG TIGTAATCAA GITTGCACAA TATATTTATC	360
55	TATATGAGGA ACCCATAAAT GAATAGCTAA TITTITAAAAT GCCATTAAAA TGCATGAAAT	420
	KCTTATTAAA ACCTTACTAT ACTATTTCTT CAAGGCAAGT AAATTGACCA TGRGRAAAGR	480
60	ACACAGITAT TAAACACTGT TGACAGGAAA ATTCTCCTTG ATAACATAGG ACAATTAATG	546

	GAAAAAAAA TICTCATTAT TIGCAAAGAA IGAACAAGIT AAIGAACAAA CAAACTAGAT	600
	TIGGTATGIT TICAGCTITT GTATCATGIT TAATIGITTA ATTIGGTIGA AAAACTGCAG	660
5	TTGAGAAATC AGATAGCAAT ATAGACATTC ACAGCAGCTC TGTGGATACC ATGTAATTGT	720
	CAGGTAATTT CAGAATGTTG AAAATTATTC AGTGCAGCCC TCATAGTATC ATACTTGAAG	780
10	AAATTGATTA CAGTTCCACT AAATTGTTGA AGATAAATTA TTTTTAAAGG TTATGAAAAC	840
10	TAAGTTATAT TAATTCATAT GTTTGATTTT TAAATCCCAC CTCCTCAAGC TATCCAATTT	900
	NCTGACTTTG AAAATAACCA TGAGAGATGC CACATTTCTC TCTGGGAAAC TACCACTCAA	960
15	AGAATAATTG TTAAAAATTA AGCTTTTAGG TATTAGAAGC TGTTATAAAG TATAAAATTA	1020
	AGATATAAGC AGATCACATG TAAATCATTC CTAAAGCACA AGAAAAGAAT GTGCCTTGAT	1080
20	GTACATATAT TACTAAGTTG CCTCTCCCAG TTTACTTTAA AAATGGCTTT AAGGATAAAG	1140
20	AATAAATGTG ATAGCTGTGC ATGCATTATA TATTTGCATT TGCAAATTTC CCATTGTTTT	1200
	AACAGCTGTG TGGCTGACTT TCAATTTTAA GACGTGAATT GACATACAGC CCATAACTTT	1260
25	ATAATGGCTG CTCATTTATC TTATCTTTCA GTTAGTGGAA AAACATTTCA ACCTGACTAA	1320
	AATTIGGAAT TGIGTCTTTT ATGITCCATC CICTGTTGTT ACTAGATTTA GITTAAAAAT	1380
30	TGTGTATGAC CATTAATGTA TGTCATAAAC ATGTAAATAA AAGATGTTGA ATCTTGTTGA	1440
	AAAGCAWRAA AAAAAAAAA AAACTCGA	1468
35	(2) INFORMATION FOR SEO ID NO: 143:	
	••	
	(i) SEQUENCE CHARACTERISTICS:	

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(A) LENGTH: 300 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 143: 45

TGAATTITTT GCCAAACTTA GTAACTCTGT TAAATATTTG GAGGATTTAA AGAACATCCC 60 AGTTTGAATT CATTTCAAAC TTTTTAAATT TTTTTGTACT ATGTTTGGTT TTATTTTCCT 120 TCTGTTAATC TTTTGTATTC RCTTATGCTC TCGTACATTG AGTACTTTTA TTCCAAAACT AGTGGGTTTT CTCTACTGGA AATTTTCAAT AAACCTGTCA TTATTGCTTA CTTTGATTAA 240 AAAAAAAAA AAAAAAAAA AAACCCCNAG GGGGGGCCG GGTNCCCAAT CCCCCCCAAA 300

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TGCCTCCCTT CCTGCAGATT GTGGACAGTA GTTCCTCAGC CTGCACCCTG GATTCCTTCT 60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2243 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 144:

10	TGCCTCCCTT CCTGCAGATT GTGGACAGTA GTTCCTCAGC CTGCACCCTG GATTCCTTCT	60
10	TCCCCTTCCT AGCTCCATGG GACTCGCCCC AAGACTGTGG CTTCAAGGAC CACCAGCCCC	120
	TTACTCTTCA AGCCCTGACT GTGGAGTTGG TAGATGCCTC TGATCCTCAG TATTCTCTCT	180
15	GCCAATGTTC CACGCCTTCT CCTTCCTGGG AGCTGGCTCC ATAACTTGAT TTTCCCCAAA	240
	CGTGTTGCAA TCCCTGCTGC CCCTTAGCCA CCCAGGGTCT TGTGTGGGTA TGAGTGTAGA	300
20	GGATGGGGGT ATGCCAGGCC TGGGCCGTCC CAGGCAGGCC CGCTGGACCC TGATGCTACT	360
20	CCTATCCACT GCCATGTACG GTGCCCATGC CCCATTGCTG GCACTGTGCC ATGTGGACGG	420
	COGAGTICCC TTYCGCCCT CCTCAGCCGT GCTGCTGACT GAGCTGACCA AGCTACTGTT	480
25	ATGCGCCTTC TCCCTTCTGG TAGGCTGGCA AGCATGGCCC CAGGGGCCCC CACCCTGGCG	540
	CCAGGCTGCT CCCTTCGCAC TATCAGCCCT GCTCTATGGC GCTAACAACA ACCTGGTGAT	600
30	CTATCITCAG CGTTACATGG ACCCCAGCAC CTACCAGGTG CTGAGTAATC TCAAGATTGG	660
30	AAGCACAGCT GTGCTCTACT GCCTCTGCCT CCGGCACCGC CTCTCTGTGC GTCAGGGGTT	720
	AGCGCTGCTG CTGCTGATGG CTGCGGGAGC CTGCTATGCA GCAGGGGGCC TTCAAGTTCC	780
35	CGGGAACACC CTTCCCAGTC CCCCTCCAGC AGCTGCTGCC AGCCCCATGC CCCTGCATAT	840
	CACTCCCCTA GECCTGCTGC TCCTCATTCT GTACTGCCTC ATCTCAGGCT TGTCGTCAGT	900
40	GTACACAGAG CTGCTCATGA AGCGACAGNG GCTGCCCCTG GCACTTCAGA ACCTCTTCCT	960
40	CTACACTTTT GGTGTGCTTC TGAATCTAGG TCTGCATGCT GGCGGCGGCT CTGGCCCAGG	1020
	SCTCCTGGAA GGTTTCTCAG GATGGGCAGC ACTCGTGGTG CTGAGCCAGG CACTAAATGG	1080
45	ACTECTCATE TCTECTETCA TEAAGCATES CASCASCATC ACACECCTCT TTETESTETC	1140
	CTGCTCGCTG GTGGTCAACG CCGTGCTCTC AGCAGTCCTG CTACGGCTGC AGCTCACAGC	1200
50	COCCTTCTTC CTGGCCACAT TGCTCATTGG CCTGGCCATG CGCCTGTACT ATGGCAGCCG	1260
50	CTAGTCCCTG ACAACTTCCA CCCTGATTCC GGACCCTGTA GATTGGGCGC CACCACCAGA	1320
	TOCCCCTCCC AGGCCTTCCT CCCTCTCCCA TCAGCAGCCC TGTAACAAGT GCCTTGTGAG	1380
55	AAAAGCTGGA GAAGTGAGGG CAGCCAGGTT ATTCTCTGGA GGTTGGTGGA TGAAGGGGTA	1440
	CCCCTAGGAG ATGIGAAGTG TGGGTTTGGT TAAGGAAATG CTTACCATCC CCCACCCCCA	1500
<i>6</i> 0	ACCAAGTTCT TCCAGACTAA AGAATTAAGG TAACATCAAT ACCTAGGCCT GAGAAATAAC	1560
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CCCATCCTTG	TTGGGCAGCT	CCCTGCTTTG	TCCTGCATGA	ACAGAGTTGA	TGAAAGTGGG	1620
GTGTGGGCAA	CAAGTGGCTT	TCCTTGCCTA	CTTTAGTCAC	CCAGCAGAGC	CACTGGAGCT	1680
GGCTAGTCCA	GCCCAGCCAT	GGTGCATGAC	TCTTCCATAA	GGGATCCTCA	CCCTTCCACT	1740
TTCATGCAAG	AAGGCCCAGT	TGCCACAGAT	TATACAACCA	TTACCCAAAC	CACTCTGACA	1800
GTCTCCTCCA	GTTCCAGCAA	TGCCTAGAGA	CATGCTCCCT	GCCCTCTCCA	CAGTGCTGCT .	1860
CCCCACACCT	AGCCTTTGTT	CTGGAAACCC	CAGAGAGGGC	TGGGCTTGAC	TCATCTCAGG	1920
GAATGTAGCC	CCTGGGCCCT	GGCTTAAGCC	GACACTCCTG	ACCTCTCTGT	TCACCCTGAG	1980
GCTGTCTTG	AAGCCCGCTA	CCCACTCTGA	GGCTCCTAGG	AGGTACCATG	CTTCCCACTC	2040
TOGGGCCTGC	CCCTGCCTAG	CAGTCTCCCA	GCTCCCAACA	GCCTGGGGAA	GCTCTGCACA	2100
GAGTGACCTG	AGACCAGGTA	CAGGAAACCT	GTAGCTCAAT	CACTGTCTCT	WTAACTGCAT	2160
AAGCAATAAG	ATCTTAATAA	AGTCTTCTAG	GCTGTAGGGT	GGTTCCTACA	ACCACAGCCA	2220
ааааааааа	AAAAAAACTC	GAG				2243

(2) INFORMATION FOR SEQ ID NO: 145:

30 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1082 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:

GCCAAGCTCT AATACGACTC ACTATAGGGA AAGCTGGTAC GCCTGCAGKT ACCGGTTCCG 60 120 GGAATTCCCG GGTCGACCCA CGCGTCCGCT TCCGTGTGTC AAAATCCTCA CCTCCTTCAT AACCATCTCC CACAATTAAT TCTTGACTAT ATAAATTTAT GGTTTGATAA TATTATCAAT 180 TTGTAATCAA TTGAGATTTC TTTAGTGCTT GCTTTTCTGT GACTCAACTG CCCAGACACC 240 TCATTGTACT TGAAAACTGG AACANCTTGG GAATGCCATG GGGTTTGATA ATCTGCCAGG 300 GACATGAAGA GOCTCAGCTT CCTGGGACCA TGACTTTGGC TCAGCTGATC CTGNACATGG 360 GAGAACAACC ACATTTTTCT TTGTGTGTGC TTCTAGCAGC TGTTCGGGAG GACCKTGACC 420 CAAYAGTGTT CCCATGCTGT TICTTGTGAA ATGCTCTCGG CTATGTAGCA GCTTTTGATT 480 CCCTGCATAC CCTAGGCTGC TGCCCCTATC CTGTCCCTTG TTTATAACAT TGAGAGGTTT 540 TCTAGGGCAC ATACTGAGTG AGAGCAGTGT TGAGAAGTCG GGGAAAATGG TGACTACTTT 600 TAGAGCAAGG CTGGGCATCA GCACCTGTCC AGCTCTACTT GTGTGATGTT TCAGGAACTC AGCCCCTTTT TCTGCCTAGG ATAAGGAGCT GAAAGATTAA CTTGGATCTY CTAATGGTCC 720

395

•	AAATCTTTTG GTCACAATAA AGAGTCTCCA AATTAGAGAC TGCATGTTAG TTCTGGATGG	780
5	ATTTGGTGGC CTGACATGAT ACCCTGCCAG CTGTGAGGGG ACCCCGTTTT TAAGATGCAT	840
)	GGCCAAGCTC TCTGCAAATG GAAATGCTTA CACTGGGTGT TGGGGATGTT TGCTACCTCC	900
	TOCTATTITT GIGGITITGG TICTCCCACT ATGGTAGGAC CCCTGGCCAG CATTGTGGCT	960
10	TGTCATGTCA GCCCCATTGA CTACCTTCTC ATGCTCTGAG GTACTACTGC CTCTGCAGCA	1020
	CAAATTTCTA TTTCTGTCAA TAAAAGGAGA TGAAAATAAA AAANAAAAAA AAAAAACTCG	1080
15	NG	1082
13		
20	(2) INFORMATION FOR SEQ ID NO: 146:	

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 4313 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

25 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:

30	CAAGCTGGTT TGAAACTAGG GGTCGGGCTC GGCCGTCGTC GTTGTTTGTC GCCGCATCCC	60
30	CGCTTCCGGG TTAGGCCGTT CCTGCCCGCC CCCTCCTCTC CTCCCTTCGG ACCCATAGAT	120
	CTCAGGCTCG GCTCCCCGCC CGCCGCAGCC CACTGTTGAC CCGGCCCGTA CTGCGGCCCC	180
35	GTGGCCACCA TGTCCCTGCA CGGCAAACGG AAGGAGATCT ACAAGTATGA AGCGCCCTGG	240
	ACAGTCTACG CGATGAACTG GAGTGTGCGG CCCGATAAGC GCTTTCGCTT GGCGCTGGGC	300
40	AGCTTCGTGG AGGAGTACAA CAACAAGGTT CAGCTTGTTG GTTTAGATGA GGAGAGTTCA	360
40	GAGTITATIT GCAGAAACAC CITTGACCAC CCATACCCCA CCACAAAGCT CATGIGGATC	420
•	CCTGACACAA AAGGCGTCTA TCCAGACCTA CTGGCAACAA GCGGTGACTA TCTCCGTGTG	480
45	TOGAGGGTTG GTGAAACAGA GACCAGGCTG GAGTGTTTGC TAAACAATAA TAAGAACTCT	540
	GATTTCTGTG CTCCCCTGAC CTCCTTTGAC TGGAATGAGG TGGATCCTTA TCTTTTAGGT	600
50	ACCTCAAGCA TTGATACGAC ATGCACCATC TGGGGGCTGG AGACAGGGCA GGTGTTAGGG	660
50	CGAGTGAATC TCGTGTCTGG CCACGTGAAG ACCCAGCTGA TCGCCCATGA CAAAGAGGTC	720
	TATGATATTG CATTTAGCCG GGCCGGGGGT GGCAGGGACA TGTTTGCCTC TGTGGGTGCT	780
55	GATGGCTCGG TGCGGATGTT TGACCTCCGC CATCTAGAAC ACAGCACCAT CATTTAGGAA	840
	GACCCACAGC ATCACCCACT GCTTCGCCTC TGCTGGAACA AGCAGGACCC TAACTACCTG	900
60	GCCACCATGG CCATGGATGG AATGGAGGTG GTGATTCTAG ATGTCCGGGT TCCTGCACAC	960

	CTGTSGCCAG GTTAAACAAC CATCGAGCAT GTGTCAATGG CATTGCTTGG GCCCCACATT .	1020
	CATCCTGCCA CATCTGCACT GCAGCGGATG ACCACCAGGC TCTCATCTGG GACATCCAGC	1080
5	AAATGCCCCG AGCCATTGAG GACCCTATCC TGGCCTACAC AGCTGNAAGG WGAGATCAAC	1140
	AATGTGCAGT GGGCATCAAC TCAGCCCGAA YTGTCGCCAT CTGCTACAAC AACTGCCTGG	1200
10	AGATACTCAG AGTGTAGTGT TGGTGGCGCT GTGCCCACGA GGCAGGGGCT TTTGTATTTC	1260
10	CTGCCTCTGC CCCACCCCCA AAGTAAGAAG AAACATGTTT CCAGTGGCCA GTATGTCTTT	1320
	CATTGCTTTG CACCCACTGT TACCAGAAGC TGCTCTAGGA GTTCCTGGCC AGTCACCCCA	1380
15	TOGOCCTOTG TOGOLAGACTO AGTOCTOTGT GOCGCCTCCT CAGCCCAGGG CTGAGTTTTA	1440
	AGATTTICTC TCCTTTCCTC TTCTCCTTTG GFTCCTCAAT TAAAAAATGT GTGTATATTT	1500
20	GTTTGTCAGG CGTTGTGTTG AGGAGCAGTT CACGCACTGG CTGTGTCTAT TCCTCTGCCC	1560
20	AGGTGTCTCT GTTTGCTGCC CAAKGYWKKT TTTCATGTCT CGTCCATGTC CATGTTCGTG	1620
	TTAGCACTWA CGTGGGAACA AATACCAATT TGTCTTTTCT CCTAGTATCA GTGTGTTTAA	1680
25	CAAATTTTAA CTTTGTTATAT TTGTTATCTA TCAGGCTAAT TTTTTTATGA AAAGAATTTT	1740
	ACTOTOCIGO TICATTICIT TOTOTIATAG TOCTOCCTOT TIGGACOTTO TICTOTICOC	1800
30	TCAGTGCCTG GAGCTGGTAC TGGGCCCCTG GCCCCATGAG CAGTTTGCCT TCTTGAGTCA	1860
50	CTGCCTGTGT AGTACATACC TGACCGGGGG TCCAAACCAC CTTGGTGCTC TGAAGTCCAC	1920
	TGACTCATCA CACCTTTCTT AGCCTGGCTC CTCTCAAGGG CATTCTGGGC TTGTAAACAG	1980
35	ACATAGGAAG CCTCTGTTTA CCCTGAAGCA CCACTGTCCA GCCCATTGGT TCCCACTGGC	2040
	AGCATGGTAG AGCTGAGAGA AACAGGCTCT CAGGGTACCT GACTTGAGGG GAATCGTTTC	2100
40	ATGAAGCTGA ACTTCAAGCA TATTTCCAGT ACATTCTTTC AGAGTCTGTT TTTCCATCCA	2160
	AATATAAGCC CCAGGCCATT CCACTTAGTG TCTTTTCAAT GATAGGCAAG AATGATATCT	2220
•	GAGTTGAACT TCGGTGCTTC TGTTGTTTGA GTTTACTGTG CCTGGTGGTA TATTGGGCAT	2280
45	TCTTTGGATT GAGTGTTCTG AGGTGAGAGA GTCTTCCCGA GGCATCCTGT CTGTGCTTCC	2340
	AACCCTGAAC AAGACCTTAC ATGAGAGATG GACTGATGGA CTGCGGCAAT CCTGGGCTGT	2400
50	CAAGTGGATA GATAGTTAAA AAGCATTATA CTGTGGGTAA TGAAAAGGGA GGAAAAAAA	2460
	AGAAGGAAAA GGAATTATAG ACCCCCAGGG TCAGCCAGTT AAGAGCTCTA CCCACACCTG	2520
	TCAACCCCTC TCTCCCCCAG TTTAGGTTCT GAGCAGTATT GGACTTGTAG CCTGCAGTTG	2580
55	TCTTTIGACT TGCAGGCCGC AGTGTCTTTC TGTTATGTGA ATGAGTTCCA TGGAGGGCA	2640
	TATGTGTGAT TCCACCGTTA GATGAGCCCT TGGGGCAGGC AGTTTGGGAT GTGCTCTTGG	270
60	GOGANAGITG GCTGTTTCCT TGCGCTCTGC TCCTACCCGA AGTTTTTANG TCCCTCTGAN	276

	TIGCTCATCT GAGATTAGTA GAGTAGCAGG CCTGAAGGAT GATGGTTTTG TCCTCTTTGG	2820
	TTCTCACCTG CTTGAGAAGT AAAACAGTAA CTTTGTTCTT CTGGGCCCTT AAGCTTTTTT	2880
5	GGTTAAGTCT TCCTTTTCAG AAGTAGATGT CATTATATGC CAAAAGTCTA GCTCTTTGCT	2940
	TTACCATACA GGGACCTGTC CCAAAGAAAA AGGCTCTTTT TTTAGCCAGC ATATTTCCCC	3000
10	TTCTACCCTT TTACTTTGTT GTTCTGATTT TAGGACTCTG GCTGGCCATG TGCTTGTGGT	3060
10	TGCCTCTCCT GCATTTGCCA CTGGATTTGC ACTGCATCGT TTGGAGATAC AAAGCGAGCA	3120
	GITCITGGIC AGAACCCICC TCTGCITTIC ATTGIGITIG ATAATGGITA CTGGGTCCTT	3180
15	CTCTCAAGGG TAGCAAGGCC AAGCTGATGG CTGCTTGTTT AGGAGGCCAT CAGTTCCTTC	3240
	CTGTGGAGAA GGGTCTGAAA TGGAAGTCAG TGGTAGAAGG GGCTGGTCTG CTGGGCAGGG	3300
20	CTTACATCCA CTGAGTTCTA AGATTCCTTT CCTGATCTGC ACCTACGCCT GGTCTGTATG	3360
20	GTGGAATTTG TCAGCTGGAA CTCAGAAACA ACAACTTGAA AAAAAAATAA TAATTAGAAC	3420
	ATATTTGCAT AAGATAGCTA TTTACTCTGG AAACCAACAA CTTTTGAGAT TTCCCTTGCC	3480
25	CTGTGGACGC CCAGCTCCTG TCATCCTTCC TTAGGTCCTG CAGTACAGTC TTCCCCTGAA	3540
	TGCCACCGGG GACCCAGGGG GACTCCACCC CCCTAAGCAA GCACACACAT ACTCACAGTT	3600
30	GATGAGTTGC TGGTCTTTGA GTCCCAGCTC TCTTACCCTC CCTTTACTCC ACCAGCCCGA	3660
	CGACCCATGA CTGAGGAGGG GATTTCTACA GTCTCAGGAT TTAGAAAGTC TGTAAGCCAT	3720
	CCATGCTCCA GAAAGCACCG ATCTGTTGTA GTTGCAAAAA CAACTCTGTA ATTTGTTGAG	3780
35	GTTCTCAAAC TGACAGCCAG CGAGACTGGG TGGGAGGCCC TGGATCTGTT CTCCCTGACT	3840
	GCGGGAGGAG CAGCCACTAG GACTITIAGCA GGAAGCCCAC ATGGAGGCTC CGCCAGGCTG	3900
40	TGGCCCAGCT GGTGATGGCC CTTTTGCTCC TGGCAGCCTG AGGCACAGCT GCCTGTATTG	3960
	TCCTCATCTG TTCTGACTGA AGGATGGAGG TGCTGAATAA ATTAGGCCTC AGGCNTCTAC	4020
	CACCAGAGAG CTGGAGAATG GGTCCACGTC ATTCAAGGAC CTGAATTTTT TATGCTCAGG	4080
45	AGCATTOGAA TCCTCTTCTT CCAGGGAGGA ATTAGCCTGC AAGGTTAGGA CTTGAAGAGG	4140
	GAAGGTATTT AATAACTGGG CGAGGATGGG TGTGGTGGCT CACACCTGTA ATCCCAGCAT	4200
50	TTTGGGAGGC TGAGGTGGCC AGATCCCAAG GTCAGAAGAT CGAGACCATC CTGGCTAACA	4260
	TOTAL A ACC CONTINUES TABABATACA ARATTARATT GGCCGGGCGT GAR	4313

(A) LENGTH: 1183 base pairs
(B) TYPE: nucleic acid

⁽²⁾ INFORMATION FOR SEQ ID NO: 147:

⁽i) SEQUENCE CHARACTERISTICS:

(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

_							
5	GGCAGAGCCT	CAAGCTGACT	TOGATTATGT	GGTCCCTCAA	ATCTACCGAC	ACATGCAGGA	60
	GGAGTTCCGG	GCCCGGTTAG	AGAGGACCAA	ATCTCAGGGT	CCCCTGACTG	TGGCTGCTTA	. 120
10	TCAKWYGGGG	AGTGTCTACT	CAGCTGCTAT	GGTCACAGCC	CTCACCCTGT	TGGCCTTCCC	180
	ACTICICCIC	TTGCATGCGG	AGCGCATCAG	CCTTGTGTTC	CTGCTTCTGT	TTCTGCAGAG	240
15	CTTCCTTCTC	CTACATCTGC	TIGCIGCIGG	GATACCCGTC	ACCACCCCTG	GTCCTTTTAC	300
13	TGTGCCATGG	CAGGCAGTCT	CGGCTTGGGC	CCTCATGGCC	ACACAGACCT	TCTACTCCAC	360
	AGGCCACCAG	CCTGTCTTTC	CAGCCATCCA	TTGGCATGCA	OCCITICATEG	GATTCCCAGA	420
20	GGGTCATGGC	TCCTGTACTT	GGCTGCCTGC	TITGCTAGTG	GGAGCCAACA	CCTTTGCCTC	480
	CCACCTCCTC	TTTGCAGTAG	GTTGCCCACT	GCTCCTGCTC	TOGCCTTTCC	TGTGTGAGAG	540
25	TCAAGGGCTG	CGGAAGAGAC	AGCAGCCCCC	AGGGAATGAA	GCTGATGCCA	GAGTCAGACC	600
	CGAGGAGGAA	GAGGAGCCAC	TGATGGAGAT	CCGCTCCGG	GATGCGCCTC	AGCACTTCTA	660
	TGCAGCACTG	CTGCAGCTGG	GCCTCAAGTA	CCTCTTTATC	CTTGGTATTC	AGATTCTGGC	720
30	CTGTGCCTTG	GCAGCCTCCA	TCCTTCGCAG	GCATCTCATG	GICTGGAAAG	TGTTTGCCCC	780
	TAAGTTCATA	TTTGAGGCTG	TOGGCTTCAT	TGTGAGCAGC	GTGGGACTTC	TCCTGGGCAT	840
35	AGCTTTGGTG	ATGAGAGTGG	ATGGTGCTGT	GAGCTCCTGG	TTCAGGCAGC	TATTTCTGGC	900
	CCAGCAGAGG	TAGCCTAGTC	TGTGATTACT	GCACTIGGC	TACAGAGAGT	GCTGGAGAAC	960
	AGTGTAGCCT	GGCCTGTACA	GGTACTGGAT	GATCTGCAAG	ACAGGCTCAG	CCATACTCTT	1020
40	ACTATCATGO	: AGCCAGGGGC	CGCTGACATO	TANGACTICA	TTATTCWATE	ATTCAGGACC	1080
	ACAGTGGAGT	* ATGATCCCTA	ACTCCTGATT	TGGATGCATC	TGAGGGACAA	GGGGGKCGGT	1140
45	STCCGAAGTC	GAATAAAATA	GCCCCCCCTC	GTGACTTGCA	CCT		1183

(2) INFORMATION FOR SEQ ID NO: 148:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 734 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

GAATTCGGCA GAGTGAAGCA TTAGAATGAT TCCAACACTG CTCTTCTGCA CCATGAGACC 6

399

	AACCCAGGGC AAGATCCCAT CCCATCACAT CAGCCTACCT CCCTCCTGGC TGCTGGCCAK	120
	GATGTCGCCA GCATTACCTT CCACTGCCTT TCTCCCTGGG AAGCAGCACA GCTGAGACTG	180
5	GGCACCAGGC CACCTCTGTT GGGACCCACA GGAAAGAGTG TGGCAGCAAC TGCMTGGCTG	240
	ACCITICIAT CITCICIAGG CICAGGIACI GCICCICCAI GCCCATGGYI GGGCCGIGGG	300
10	GAGAAGAAGC TCTCATACGC CTTCCCACTC CCTCTGGTTT ATAGGACTTC ACTCCCTAGC	360
10	CAACAGGAGA GGAGGCCTCC TGGGGTTTCC CCRRGGCAGT AGGTCAAACG ACCTCATCAC	420
	AGTOTTCCTT COTOTTCAAG COTTTCATGT TGAACACAGC TCTCTCCRCT CCCTTGTGAT	480
15	TTCTGAGGGT CACCACTGCC ARCCTCAGGC AACATAGAGA GCCTCCTGTT CTTTCTATGC	540
	TTGGTCTGAC TGAGCCTAAA GTTGAGAAAA TGGGTGCCAA GGCCAGTGCC AGTGTCTTGG	600
20	GGCCCCTTTG GCTCTCCCTC ACTCTCTGAG GCTCCAGCTG GTCCTGGGAC ATGCAGCCAG	660
20	GACTGTGAGT CTGGGCASGT CCAAGGCCTG CACCTTCAAG AAGTGGAATA AATGTGGCCT	720
	TIGCTICTAT TTAA	734
25		
	(2) INFORMATION FOR SEQ ID NO: 149:	
30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1405 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 149:	
	GGCACAGTGG ACCCCAGACT CCCTCTCCGC CTTTCTCTGC CTGGGGAGAC CCACTGTGTG	60
40	CATGGCATCA CTGACTCCCA TACCTCTGGC TATCAAAGGT TTCTGCCATG GCCACCCTGG	120
	AAGSAAACCA GAGGGAGGTA GACAGGGAGA TCAGGTCCCT TCTACTCTGG TTCCTGCTCT	180
	GTGAAATTGT CTCAGGCTGG CTGTGTCCAG ARGGTCCCTG GTTCTCTCAR GGATGCCAAA	240
45	TCTACAAGAA TCTCTCCTCT TCCAGTTCCT ATAACCTCTC CTTCCTTTTG TCTCTTTAGA	300
	CCTTGGAGTA GTAGCAGCCA GGTTCTTTCT ATCTCTGGGT TAGTGCATTA TCTCTGGTGG	360
50	CTCCCTTACC CAGGACTITG GGAATGGTCT TTTTGTAATA CATTCTCCTC AAATAATTCA	420
	ATTTGAGIG TTCTGTATGT ATCCTGCTGG GAGGTTGTTA TATACAAATC ACTGTGCCCG	480
e e	TITAGCAGAG AAGGAGACTG AAGCTCAGGG AGGTTAAGTG TCTTTCTCTA GGTCGTATTG	540
55	TGGAGAAAGT GGCTGACTGG GGACTTGAAT GAGGTCCCTA GTTTCATGCT CGGAGGGCAA	600
	AGANGAATGT CCAATTOGCC TGAGATAAGC CTCTGGTAAA ATGTACTGTA CATAATAGGT	660

AATCAATAAA TGTTGGCTGA TGACAAACAT GTTTTCTTTG TTCATTAGTT ATAGTGATTA

60

400

	TGTTCTAAAT AACTCCMACA AGGAARTCAG CACATTTGGA ATATCAWTAT CTTTCCATGA	780
5	TAATATCTTT COMYGGAAAG AWAATGATAT TOOMAACTGG GAGTGTCCCW AGCARATCTG	840
J	ANICIGISTA TIGGCCCIGG GGIGGGCCAG CCCCTTAGAC TCTATGGTCT CATTCTCTTT	900
	GTTTACAAAA TTGAGATAAG GCCTTATTCT CTCCCCACCC CACCCATCCA TATTGTTTTG	. 9 60
10	AGAATAAAAT GAGAGGATGT GTGTCAAGGG TGTATTTTGG CAATAGTCTC TGAGCCATTT	1020
	TCTGAGCACC TCCATACTGT TGACACTCAA GTAATATTTC ATCAGCATTC CATTCAGGNT	1080
15	CCTCCCTTAA TGAGGTGTGC GATGTACAAG AGTYGTGAGG TGGCAAAGGA TGGGCTCCTG	1140
13	AGGAAACACT TAGGAAACTG GGCTTTCTGC CATTAAAAGA GACAAACCTT TGTGGTGACC	1200
	TAATTAAAGT TITTAAAATT CAATTIGGAA AGITAGCAAG CTAGCTCCTK TCCAGGWAAA	1260
20	ATAAGGAGTC AGTGCATGAC CTAACCGGTC CCGGGCTGCT TGCCATTCCA AACAACTGCA	1320
	GTAAGTTTAT CACNITCTIT CAGGGACTGA GGTTTCCAGG CACAGACTTG GATAAGGAAG	1380
25	GATGTCCTAT GGGGTCACAT TGATG	1409
23		

(2) INFORMATION FOR SEQ ID NO: 150:

30

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2890 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 150:

40	TTATATGCTA CAGCTACAGT AATTTCTTCT CCAAGCACAG AGGANCTTTC CCAGGATCAG	60
40	GGGGATCGCG CGTCACTTGA TGCTGCTGAC AGTGGTCGTG GGAGCTGGAC GTCATGCTCA	120
	AGTGGCTCCC ATGATAATAT ACAGACGATC CAGCACCAGA GAAGCTGGGA GACTCTTCCA	180
45	TTCGGGCATA CTCACTTTGA TTATTCAGGG GATCCTGCAG GTTTATGGGC ATCAAGCAGC	240
	CATATOGACC AAATTATGTT TTCTGATCAT AGCACAAAGT ATAACAGGCA AAATCAAAGT	300
50	AGAGAGAGCC TTGAACAAGC CCAGTCCCGA GCAAGCTGGG CGTCTTCCAC AGGTTACTGG	360
30	GGAGAAGACT CAGAAGGTGA CACAGGCACA ATAAAGCGGA GGGGTGGAAA GGATGTTTCC	420
	ATTGAAGCCG AAAGCAGTAG CCTAACGTCT GTGACTACGG AAGAAACCAA GCCTGTCCCC	480
55	ATGCCTGCCC ACATAGCTGT GGCATCAAGT ACTACAAAGG GGCTCATTGC ACGAAAGGAG	540
	GGCAGGTATC GAGAGCCCCC GCCCACCCCT CCCGGCTACA TIGGAATTCC CATTACTGAC	600
60	TTTCCAGAAG GGCACTCCCA TCCAGCCAGG AAACCGCCGG ACTACAACGT GGCCCTTCAG	660

	AGATCGCGGA TGGTCGCACG ATCCTCCGAC ACAGCTGGGC CTTCATCCGT ACAGCAGCCA	720
	CATGGGCATC CCACCAGCAG CAGGCCTGTG AACAAACCTC AGTGGCATAA AYCGAACGAG	780
5	TCTGACCCGC GCCTCGCCCC YTATCAGTCC CAAGGGTTTT CCACCGAGGA GGATGAAGAT	840 -
	GAACAAGTTT CTGCTGTTTG AGGCACAGAC TTTTCTGGAA GCAGAGCGAG CCACCTGAAA	900
	GGAGAGCACA AGAAGACGTC CTGAGCATTG GAGCCTTGGA ACTCACATTC TGAGGACGGT	960
10	GGACCAGITT GCCTCCTTCC CTGCCTTAAA AGCAGCATGG GGSTTCTTCT CCCCTTCTTC	1020
	CTTTCCCCTT TGCATGTGAA ATACTGTGAA GAAATTGCCC TGGCACTTTT CAGACTTTGT	1080
15	TOCTTGAAAT GCACAGTGCA GCAATCTTCG AGCTCCCACT GTTGCTGCCT GCCACATCAC	1140
	ACAGTATCAT TCCAAATTCC AAGATCATCA CAACAAGATG ATTCACTCTG GCTGCACTTC	1200
20	TCAATGCCTG GAAGGATTTT TTTTAATCTT CCTTTTAGAT TTCAATCCAG TCCTAGCACT	1260
20	TGATCTCATT GGGATAATGA GAAAAGCTAG CCATTGAACT ACTTGGGGCC TTTAACCCAC	1320
	CAAGGAAGAC AAAGAAAAAC AATGAAATCC TTTGAGTACA GTGCTTGTCC ACTTGTTTAC	1380
25	AATGTCCTCC TTTTAAAAAA AAAAAAATGA GTTTAAAGAT TTTGTTCAGA GAGTAAATAT	1440
	ATATCCATTT AATGATTACA GTATTATTTT AAACCTTAAG TAGGGTTGCC AGCCTGGTTT	1500
30	CTGAAAAACC AAATATGCCG GACAGGGTGT GGCCACACCA AGAAGACGGG AAGACCTGGC	1560
50	TIGIGACCCT GECTICCCAT GICCTICIGG TCTCACCCGC GAAGIGCCCT ATCCTGGAAG	1620
	TATGAAATGT TAGCCAATTA ATACCAAGAC ACCTCATCTG CTCCTTCCCC AGTGGATGGG	1680
35	GTICTICIGI AAAACIGITT GCACATGGCC AGGGGAGGGA ACTAGGACCC TIGTGTCCIG	1740
	TCTGAGCCTT ATGGAGGCAG GACQGTGTCA TTGGCGGATG TGTCCTGCTC CATTGAGATG	1800
40	GATGGCAAAC CCCATTTTTA AGTTATATTT CTTTGATTTT TGTTAATTTA GAGGTGTAGG	1860
40	TTTTGTTTTT TGTTTTTTG TTTTTTTTTA AGAGAAACAT TTATAACTGG ATAGCATTGC	1920
	AGTGAAAGCA GCTTGGGATG TTGGAGCTAA TGCCAGCTGT TTATACTGCT CTTTCAAGAC	1980
45	AGCCTCCCTT TATTGAATTG GCATTAGGGA ATAAACAAGC CTTTAAACGT GATAAAAGAT	2040
	CAAAAACCTG GTTAGACATG CCAGCCTTTG CAAGGCAGGT TAGTCACCAA AGACTAACCT	2100
50	CCAAGTGGCT TTATGGACGC TGCATATAGA GAAGGCCTAA GTGTAGCAAC CATCTGCTCA	2160
30	CASCISCIAT TAACCCIATA ATGACIGAAA IGACCCCICC ACTCIATITI IGIGITGITI	2220
	TOCACAGACT CCGGAAAAGT GAAGGCTGCC AATCTGAGTA GTACTCAAAT GTGAGGAACT	2280
55	GCTGGTCTTG GATTTTTTT CCATTAAATT CAGCTGATCA TATTGATCAG TAGATAAACG	2340
	TARATAGCTT CARATTTAR RAGIGGRATT GCAGIGITTT TICACTGIRT CARACRATGI	2400
60	CAGTOCTTTA TITAATAATT CTCTTCTGTA TCATGGCATT TGTCTACTTG CTTATTACAT	2460

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	TGTCAATTAT GCATTTGT	A TTTTACATGT	AATATGCATT	ATTTGCCAGT	TTTATTATAT	2520
	AGGCTATGGA CCTCATGTO	C ATATAGAAAG	ACAGAAATCT	AGCTCTACCA	CAAGTTGCAC	2580
5	AAATGITATC TAAGCATT	a gtaattgtag	AACATAGGAC	TGCTAATCTC	AGTTCGCTCT	2640
	GTGATGTCAA GTGCAGAA	rg tacaattaac	TGGTGATTTC	CTCATACTTT	TGATACTACT	2700
10	TGTACCTGTA TGTCTTTT	ag aaagacattg	GTGGAGTCTG	TATCCCTTTT	GIATTTTTAA .	2760
10	TACAATAATT GTACATAT	rg gytatatytt	TGTTGAAGAT	GGTAGAAATG	TACTATGTTT	2820
	ATGCTTCTAC ATCCAGTT	TG TACAAGCTGG	, алаатааата	AATATAACAT	АААААААА	2880
15	АААААААА					2890

20 (2) INFORMATION FOR SEQ ID NO: 151:

25

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2399 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 151:

30	GAACTITICC ATCIGGCAAA COGGAAACTC CATCCCCATT AAACCAACTC CCCCTTTIGG	60
	TTTCCCCCCC AGNOGAATAG AATTTGGACN CCCATATAAA TCCAGGAAAC CACCTAAATT	120
25	CTTTAGTNGT TTGTGTTTGC AAGATCTAAG GTCATGGTAA ACATTAAGTT CTTAAAATTT	180
35	TTGGGAGGGA CCAGTGCACC TCTCCCTCTG AATTGTTCNC CAATTTAAAA TTGGAGTAAG	240
	GTTTTAAAAT GTCTNATTCC ATTGGAAGGG TNTGTTATTT CATTTTGAGC CCAGAGGGGA	300
40	GAGGCACATT TTAAATATCA GAATTAGATT AGCTTTGAGT TTGTACAATT GGGAACATAA	360
	TAGATTTICA TAAATTATGT GIGCCTTGTT GGAAGTGICA ACTGTCTTTA TGTCTGCTTG	420
4 5	TARAAGTTTC ARAATATGTT TTCCCTCARA AAGGCARCGT TACTTCATTT GCTTGARTAT	480
45	TATGATAGGA ATGCTTACTG ATATTACTTG ATAGTCATAT ATAGCCTAGG AAATTTAACA	540
	TATATATAAC TATAGCAGTA TTAATAATGA TAGTTGTACT TCTTTAAAAC ATTAAATTTG	600
50	AGGARACTIT RATGCTGTCT CGTGTACATT GCTTTACTAC AGTGAGGGGG AATATCCTTT	660
	AGATTGAGCC TCAATTTACT GGTTAGTAGT ATGTGAACTC TGGTATAAAA ACGTAAACTA	720
	GACAGTAGAG CCGATGAATT AAAATTGTAA ATTGCTACAT TGGCATTTTC TACCTCCTTT	780
. 55	TOTGICAGAG TATTACTITI TOCAGCATTI ATTOTTATIT GIGAGIAAAG AGGAAATGGG	840
	AACCTGAGGT TAAAATTGAC ATTTTTGTTT CATTGAGAAT TTAAGCAGTA GGTACAGGAG	900
60	AAGTGACTTG TCACATTAAT TTGGTGCCTA AATCTGTAAC TACAAGTTGT GATCGACATG	960

	TACAAAATGT CTAAGAAAGG TCATATGCTG AATATTITAC TTTTCCTGTA TAGTCTGCAT	1020
-	GATTTGTTTC ATAAACCCAG CTTATTTCCT CCAAAAAGCA AAATGGTCCT GTAATTTTTA	1080
5	AAGTAAAATA AACGIGCCAT TITGICTGCA ATCTATAATT TCAGGAAGTT ATTGRAAGTT	1140
	CTGACTCAGG GCTTTTTAAC AGTTCAAGCA ATTGTCAGTT ATATTTTGGA AACTCCATCT	1200
10	GTGTAATTCT CCAGTGCCTT GAAAGAATTA TTAACTTGGC AACACTATTA AAACTTTATA	1260
	AAAGATGGTC TTTAGTGCAC GTGTATCATT ATATACACGT TTTAAAGTCA TATTGCTTAG	1320
15	CTTGTTAATA ATGATTCTGC ATGTGTGCTG GGTTTGGGTA ATTCTTTAAA GGAAGTTTTC	1380
13	TAGATTTGCA CTTGATGTTT GTTTTTTAAA AACTGATTAT TTATGGCCGT GACACTGTTA	1440
	CCAGAAAAGT AATTCTAATT AAGTTATTAT GCAAAGTCAT CTATAAGTAG CATCTGGGAA	1500
20	GAGGAGATSG AGGCCACAGT TTGCTATTTT AGTATGAAAG GAGGATCTGT TTGGGAAACA	1560
	TAGATTGTCT TCCCCTCAAA TGAGGGGAAA AAAAAAGACC CTTTGTTCAA ATGGATTCTG	1620
. 25	TTGTAAAAAA TTATTTTAA AGGAAATCAC AAATTGTATG TCATTCTTAA TGCTAGTCTT	1680
. 23	ATAGAATAAA TCCATAAAAT TGTTTTTATG TTCAGTATGT TTATGTCATT CTAAATGCAG	1740
	CARATTCART GATAGCAGTT CARTIGACTC ATAGCAGTGT TITGTATTTT TICTARTTCT	1800
30	TTAGCTITCA ATATTGGATT AAAGTCTTGT TTGTGAATAT AGTTTCCGTA TGGCAAATGA	1860
	TTTCTTGCTT ATTAGCTTTT GTTAAAGAAT GCTTAGTAAG AGCTAAGCTT TTAAAAGTAA	1920
35	TECAAACATT TATCETTAAT AAAACCTATE GIGTAATATC ATATAATECT TTTCTTTEAT	1980
	CTTTGGAGAA TTATTCTTTT ÅTAGTAGTAT ACATGAATTT TGATTTTTAA AGCATTTAAA	2040
	AACAAATCTC AATACATTAA AAAACCTGTT ATTGTTAAAA RGGAAATTAC CATGCCTTTA	2100
40	AGAAACAAGG ATGTACATCT TCAATTCAGC ATRAGTGTCC ACATCTAGAA GGCTCTCATT	2160
	GCAGTTGTTT ACAGTTAAGG TACCTCTATC TAAAGGGCCA AAGAAGCATT TCATAYTTTA	2220
45	ACACCTCACA TICTITCAGG ATTAAGACAT ATGAAAATAG TCTGAATAGG ATAAATITGG	2280
	ATAGGAAGTA ACTTAACCAG TCTGGGAAGA TTCAGGCTTT TTCTATKAAA AAGCTTATTC	2340
	CTCTTCACAA CTCTCCCT ACCINTTCAT TTTTCAAGAG GGTAGATATT TTAAAGCCA	239

(2) INFORMATION FOR SEQ ID NO: 152:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 802 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 152:	
	CGTGCCTGTA GTAAGCTCAT CCCTGCCTTT GAGATGGTGA TGCGTGCCAA GGACAATGTT	60
5	TACCACCTGG ACTGCTTTGC ATGTCAGCTT TGTAATCAGA GATTNTGTGT TGGAGACAAA	120
	TTTTTCCTAA AGAATAACWT GAYCCTTTGC CARACGGACT ACGAGGAAGG TTTAATGAAA	180
10 -	GAAGGTTATG CACCCCMGGT TCGCTGATCT ATCAACATCA CCCCATTAAG AATACAAAGC	240
10	ACTACATICT TITATCTTTT TIGCTCCACA TGTACATAAG AATTGACACA GGAACCTACT	300
	GAATAGCGTA GATATAGGAA GOCAGGATGG TTATATGGAA TAAAAGGCGG ACTGCATCTG	360
15	TATGTAGTGA AATTGCCCCA GTTCAGAGTT GAATGTTTAT TATTAAAGAA AAAAGTAATG	420
	TACATATGGC TGGATTTTTT TGCTTGCTAT TCGTTTTTGT GTCACTTGGC ATGAGATGTT	480
20	TATTITGGAC TATTGTATAT AATGTATTGT AATATTTGAA GCACAAATGT AATACAGTTT	540
20	TATTGTGTTA CCATTTGTGT TCCATTTGCT YCTTTGTATT GTTGCATTTA GTACAATCAG	600
	TGTTTAAACT TACTGTATAT TTATGCTTTC TGTATTTACC AGCTATTTTA AATGAGCTGT	660
25	AACTITCTAG TAAAGAATTG AAAAGCAAAT CCTCACTAAA GGATACACAG GATAGGATAA	720
	AGCCAAGTON CATCAACATT AAAAAATACT AAAANANAAA ACACAAAAAA AAAAAANCCC	780
30	GGGGGGGCC CGGAACCCAT TC	802
	(2) INFORMATION FOR SEQ ID NO: 153:	
35	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH:-461 base pairs (B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 153:	
	CTAGGAGCAC CGAGCAGCTT GGCTAÀAAGT AAGGGTGTCG TGCTGATGGC CCTGTGCGCA	60
45	CTGACCOGCG CTCTGCNCTC TCTGAACCTG GCGCCCCGGA CCGTCGCCGC CCCTGCCCCG	120
	AGTOTISTICO COGCOGOCCA GATGATGAAC AATGGCOTOC TOCAACAGOO CTOTIGOOTTG	180
50	ATGITGCTCC CCTGCCGCCC AGITCTTACT TCTGTGGCCC TTAATGCCAA CTTTGTGTCC	240
	TGGAAGAGTC GTACCAAGTA CACCATTACA CCAGTGAAGA TGAGGAAGTC TGGGGCCCGA	300
	GACCACACAG GTGGGAACAA GGACAGGGGG ATTTAAGCAG TCAAAAGGAA AAACATGTTA	360
55	AGACCCTAGA CTTGTATATT GACACACTTG TACCTTGTAA GGCAGAGGAA TGTAATTAAA	420
	AAGCACTTAT TTGGCWNAAA AAAAAAAAAA AAAAAAAAA C	46

1	21	INFORMATION	FOR	SEO	ID	NO:	154:
٦		THEOMETICA	LOL	يحد			

5 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2388 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 154:

GCCCACGCGT CCGAAAGCGG AGAACGCTGG TGGGCCTGTT GTGGAGTACG CTTTGGACTG 60 AGAAGCATCG AGGCTATAGG ACGCAGCTGT TGCCATGACG GCCCAGGGGG GCTGGTGGCT 15 120 180 AACCGAGGCC GGCGCTTCAA GTGGGCCATT GAGCTAAGCG GGCCTGGAGG AGGCAGCAGG GGTCGAAGTG ACCGGGGCAG TGGCCAGGGA GACTCGCTCT ACCCAGTCGG TTACTTGGAC 240 20 300 AAGCAAGTGC CTGATACCAG CGTGCAAGAG ACAGACCGGA TCCTGGTGGA GAAGCGCTGC TGGGACATCG CCTTGGGTCC CCTCAAACAG ATTCCCATGA ATCTCTTCAT CATGTACATG 360 GCAGGCAATA CTATCTCCAT CTTCCCTACT ATGATGGTGT GTATGATGGC CTGGCGACCC 25 ATTCAGGCAC TTATGGCCAT TTCAGCCACT TTCAAGATGT TAGAAAGTTC AAGCCAGAAG 480 TITCTTCAGG GTTTGGTCTA TCTCATTGGG AACCTGATGG GTTTGGCATT GGCTGTTTAC 540 30 AAGTGCCAGT CCATGGGACT GTTACCTACA CATGCATCGG ATTGGTTAGC CTTCATTGAG CCCCCTGAGA GAATGGAGTT CAGTGGTGGA GGACTGCTTT TGTGAACATG AGAAAGCAGC 660 GCCTGGTCCC TATGTATTTG GGTCTTATTT ACATCCTTCT TTAAGCCCAG TGGCTCCTCA 35 GCATACTCTT AAACTAATCA CTTATGTTAA AAAGAACCAA AAGACTCTTT TCTCCATGGT 780 GGGGTGACAG GTCCTAGAAG GACAATGTGC ATATTACGAC AAACACAAAG AAACTATACC 40 ATAACCCAAG GCTGAAAATA ATGTAGAAAA CTTTATTTTT GTTTCCAGTA CAGAGCAAAA 900 CAACAACAAA AAAACATAAC TATGTAAACA AGAGAATAAC TGCTGCTAAA TCAAGAACTG 960 1020 TIGCAGCATC TCCTTTCAAT AAATTAAATG GTTGAGAACA ATGCATAAAA AAAGTTGCAC 45 AAGTTCCTTA TTTTCCTTAA TATTTCACTT CTATTTAATA CAAGCTGGGA CATAAAAATT 1080 1140 CTGTTGGGGA TACCTGGGGG AAGATGTGAG AAACTAATGC TGAATTCAGC TTATACATGA 50 TGAAAAGAAA AACCAGACAA AAGGAGCACA TAAATATGCA TACAGTGTAA CTGTTATTAT 1200 TTTAATACCC ACGATAAGGG ATTTTTGTTA GCATGTTTAG GGGGAACGAG GATTGGTGGG 1260 55 ATCCTTGGGG CCACAGGAAT CTGAGGCAAC GGAAGATATA TAGAGTGATC GTCCCCCTGC 1320 CGAAGGAACC TGGCAYCTGT CAAGCAGATG CTGCAGTTCA AACTTCAGCT TTTAAGATAG 1380 1440 ATAGCTATTG AAGGCAGAGG GTCAGCAGGA GGATGTGTAT TTCTAATCTA CCCTGGTAAA

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	GTCATAGGTA AGACTCAAAA GCGGGATCTT ATTCAAAAGG CAGGTATTTC CTTTGTTTTC	1500
	TGTCTTGAAA TAGCCCCTTC CCCTAAGGTG CATTCTCTCA AGTTTTCAGT ATTGCTTTAT	1560
5	TTGCAGTGAT TAAAAGAGAT GAGAGACTTT GGAGACAGAC AACGTAAGCA ACACATACAC	1620
	ACATGAAATA CTCTAGACAG AGATGAATAT AAATCTGGCC TAATAACCAG TTTTCCATGT	1680
10	AACAGTGATT TIGTGTTTCG GGCTGAAGCA GTGGTTATAT TAAAAGCCAC TAATTCCCTT	1740
10	ATCCCTTTAA AAGATTTTTA CAATTCTCCA ACCACAAACA GCACTTCTAA AACTAACTTT	1800
	ACTITICTICC CATAATITIST TOTACATIGGA AAAAAAAAAT ATTACTITIGG CCAGGGGTGT	1860
15	GTGTAAATGT GGCAGAATTC CTAGGCAGGC TGACCTTTAC AGTATGGGCC TTTAAGATAC	1920
	TOGATCCTOG TTGGGCAACA AGTGTCACGC CTGAAGTTTC TGAAAACAAA TTAGAAGACT	1980
20	GITGGCTIGG CTAATCICGT AGTICAGGGC CAAGITICIG TAGTCAGAAT GAAGAATAAA	2040
20	ATTGAAAGAA AAAGGGGGAA ATGCTTATAC TTGGCATTAA GTTGAATGCC TCAAGTCTTA	2100
	ACTATGCTT TGTAGATGAG GCAAAAGATT TCTTAGTGGT AAAATTTCTT CAACAGGTCA	2160
25	ATGCCAATCT GTATGCCATT TTAGTAAAGT AGGTAAGGAG AGTAGCCGCT CAGTAACTIT	2220
	GGCACTAAAG AAAGAGTGTG GCTCTAGAAC TTCCAATCCC ATTGCTAGAT GTGCCCTTTA	2280
30	AAAGATGGTC CAGTGCTTTC AGGGAAGGAT GTTTAGCCAG TTTTCCTAGT ATTTGTTCCT	2340
30	TAAGATTTTT TGACCTGTGC TTAATAAGAC GGACGCGTGG GTCGACCC	2388
35	(2) INFORMATION FOR SEQ ID NO: 155:	
	AT A CONTRACTOR OF A DROTTED TOTAL CO.	

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 642 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

NEE DESCRIPTION: SEO ID NO: 155:

45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 155:	
45	AAAACAGACC ATTTAAAAAC TCAGACAAGA TTATATTTAA TATATTAATT ACTAAAAAGG	60
	CACAAGATTA CACTGAACAT ATTAGCTACT AAAAAGGCAC TGCTAAGACA TTCAAGCAAA	120
50	TAGCTATTAC ACACTACTGC AGATTTTACA GGTTTCTAAT TCTAACATAT GTTTGAAAAA	180
	TCCGTGAGTA TTCCAAAATA TATTTAATAA TGGAATATCT GCATTAATAT ACCATCCATG	240
55	TGTTTTTACC ATTTGCCTTA ATATTGAATA TACTGTTTAC CTCACACTAA AAAGAAAACC	300
	AGAAGCCITA TITGTGATTT TOGGAGTOGA AGCTTCCATT TITGTGTCAA AAATGAATCC	360
	TGATTCTTAT GGAAATCTCT GTTATTAAGA TATTTCAAGA TGAGACAACA CTGAAGATCA	.420
60	AATTGTGTTT AGTATCACTA TCTTCTCTCC TCGTTTCTCT CTTACTCCTC ATCCTCCCAG	480

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	AATCTACCAG	TTTATGGTAG	AAAGATOOGA	ACCTTATTTG	AATGTGTTTT	TTTTTTCCA	54
5	TGATGTCCAA	TTTTGTTGTG	GGAAAGGATT	TOGATAAAAT	TTTTGTTTAA	ATTTTGGTAG	60
J	ATTTTTATCT	ATACAAATTT	TTAAAATTAAA	ATGTTTTGTA	AG		64
10						•	
	(2) INFORMATION FOR SEQ ID NO: 156:						

(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 1251 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 156:

20 GCCGCTGCCC CTCCACGGAG TTGCTGATCA TCTGGGCTGT GATCCACAAA CCCGGTTCTT 60 TGTCCCTCCT AATATCAAAC AGTGGATTGC CTTGCTGCAG AGGGGAAACT GCACGTTTAA 120 25 AGAGAAAATA TCACGGCCG CTTTCCACAA TGCAGTTGCT GTAGTCATCT ACAATAATAA 180 ATCCAAAGAG GAGCCAGTTA CCATGACTCA TCCAGGCACT GAGCATATTA TTGCTGTCAT 240 GATAACAGAA TTGAGGGGTA AGGATATTTT GAGTTATCTG GAGAAAAACA TCTCTGTACA 30 AATGACAATA GCTGTTGGAA CTCGAATGCC ACCGAAGAAC TTCAGCCGTG GCTCTCTAGT 360 CTTCGTGTCA ATATCCTTTA TTGTTTTGAT GATTATTTCT TCAGCATGGC TCATATTCTA 420 35 CTTCATTCAG AAGATCAGGT ACACAAATGC ACGCGACAGG AACCAGCGTC GTCTCGGAGA 480 TGCAGCCAAG AAAGCCATCA GTAAATTGAC AACCAGGACA GTAAAGAAGG GTGACAAGGA 540 AACTGACCCA GACTTTGATC ATTGTGCAGT CTGCATAGAG AGCTATAAGC AGAATGATGT 600 40 CGTCCGAATT CTCCCCTGCA AGCATGTTTT CCACAAATCC TGCGTGGATC CCTGGCTTAG 660 TGAACATTGT ACCTGTCCTA TGTGCAAACT TAATATATTG AAGGCCCTGG GAATTGTGCC 720 45 GAATITGCCA TGTACTGATA ACGTAGCATT CGATATGGAA AGGCTCACCA GAACCCAAGC 780 TGTTAACCGA AGATCAGCCC TCGGCGACCT CGCCGGCGAC AACTCCCTTG GCCTTGAGCC ACTTCGAACT TCGGGGATCT CACCTCTTCC TCAGGATGGG GAGCTCACTC CGAGAACAGG 900 50 AGAAATCAAC ATTGCAGTAA CAAAAGAATG GTTTATTATT GCCAGTTTTG GCCTCCTCAG 960 TGCCCTCACA CTCTGCTACA TGATCATCAG AGCCACAGCT AGCTTGAATG CTAATGAGGT 1020 55 AGAATGGTTT TGAAGAAGAA AAAACCTGCT TTCTGACTGA TTTTGCCTTG AAGGAAAAAA 1080 GAACCTATTT TTGTGCATCA TTTACCAATC ATGCCACACA AGCATTTATT TTTAGTACAT 1140 TTTATTTTTT CATAAAATTG CTAATGCCAA AGCTTTGTAT TAAAAGAAAT AAATAATAAA 1200 60

408

ATAAAAAAA AAAAACCCCG GGGGGGCCCC GGTCCCCAAT TGGCCCTATG G

5

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(2) INFORMATION FOR SEQ ID NO: 157:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2127 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 157:

15 CCGGCGGGAG AGGGAAGCTG CAGCGAGAGG CGCGGATCTC AGCGCGGGAG CAGTGCTTCT 60 GCGGCAGGCC CCTGAGGGAG GGAGCTGTCA GCCAGGGAAA ACCGAGAACA CCATCACCAT 120 GACAACCAGT CACCAGCCTC AGGACAGATA CAAAGCTGTC TGGCTTATCT TCTTCATGCT 180 20 GGGTCTGGGA ACGCTGCTCC CGTGGAATTT TTTCATGACG GCCACTCAGT ATTTCACAAA 240 300 CCGCCTGGAC ATGTCCCAGA ATGTGTCCTT GGTCACTGCT GAACTGAGCA AGGACGCCCA 25 GGCGTCAGCG CNCCCTGCAG CACCCTTGCC TGAGCGGAAC TCTCTCAGTG CCATCTTCAA 360 420 CAATGTCATG ACCCTATGTG CCATGCTGCC CCTGCTGTTA TTCACCTACC TCAACTCCTT CCTGCATCAG AGGATCCCCC AGTCCGTACG GATCCTGGGC AGCCTGGTGG CCATCCTGCT 480 30 GGTGTTTCTG ATCACTGCCA TCCTGGTGAA GGTGCAGCTG GATGCTCTGC CCTTCTTTGT 600 CATCACCATG ATCAAGATCG TGCTCATTAA TTCATTTGGT GCCATCCTGC AGGGCAGCCT 35 GTTIGGTCTG GCTGGCCTTC TGCCTGCCAG CTRACACGGC CCCCATCATG AGTGGCCAGG GCCTAGCAGG CTTCTTTGCC TCCGTCGCCA TGATCTGCGC TATTGCCAGT GGCTCGGAGC 720 TATCAGAAAG TGCCTTCGGC TACTTTATCA CAGCCTGTGC TGTKATCATT TTGACCATCA 780 40 TCTGTTACCT GGGCCTGCCC CGCCTGGAAT TCTACCGCTA CTACCAGCAG CTCAAGCTTG 840 AAGGACCCGG GGAGCAGGAG ACCAAGTTGG ACCTCATTAG CAAAGGAGAG GAGCCAAGAG 900 45 960 CAGGCAAAGA GGAATCTGGA GTTTCAGTCT CCAACTCTCA GCCCACCAAT GAAAGCCACT CTATCAAAGC CATCCTGAAA AATATCTCAG TCCTGGCTTT CTCTGTCTGC TTCATCTTCA 1020 CTATCACCAT TGGGATGTTT CCAGCCGTGA CTGTTGAGGT CAAGTCCAGC ATCGCAGGCA 1080 50 GCAGCACCTG GGAACGTTAC TTCATTCCTG TGTCCTGTTT CTTGACTTTC AATATCTTTG 1140 ACTOGTTGGG CCGGAGCCTC ACAGCTGTAT TCATGTGGCC TGGGAAGGAC AGCCGCTGGC 1200 55 TGCCAAGCTG GNTGCTGGCC CGGCTGGTGT TTGTGCCACT GCTGCTGCTG TGCAACATTA 1260 ACCCCCCCC CTACCTGACT GTGGTCTTCG ACCACGATGC CTGGTTCATC TTCTTCATGG 1320 CTGCCTTTGC CTTCTCCAAC GGCTACCTCG CCAGCCTCTG CATGTGCTTC GGGCCCAAGA 1380 60

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	AAGTGAAGCC AGCTGAGGCA GAGACCGCAG AGCCATCATG GCCTTCTTCC TGTGTCTGGG	1440
_	TOTGGCACTG GGGGCTGTTT TOTCCTTCCT GTTCCGGGCA ATTGTGTGAC AAAGGATGGA	1500
5	CAGAAGGACT GCCTGCCTCC CTCCCTGTCT GCCTCCTGCC CCTTCCTT	1560
	ATCCTGAGTG GTCTGGCGGT TTTTTCTTCT AACTGACTTC TGCTTTCCAC GGCGTGTGCT	. 1620
10	GGGCCCGGAT CTCCAGGCCC TGGGGAGGGA GCCTCTGGAC GGACAGTGGG GACATTGTGG	1680
	GTTTGGGGCT CAGAGTCGAG GGACGGGTG TAGCCTCGGC ATTTGCTTGA GTTTCTCCAC	1740
15	TCTTGGCTCT GACTGATCCC TGCTTGTGCA GGCCAGTGGA GGCTCTTGGG CTTGGAGAAC	1800
13	ACGIGIGICT CIGIGIATGI GICTGIGIGI CIGCGICCGI GICTGICAGA CIGICIGCCI	1860
	GTCCTGGGGT GGCTAGGAGC TGGGTCTGAC CGTTGTATGG TTTGACCTGA TATACTCCAT	1920
20	TETECCETGE GESTECTECT STGTGTTSTC TECATGTCCC SCTCCCAACT CCCCATGCCC	1986
	AGITCITACC CATCATGCAC CCTGTACAGT TGCCACGTTA CTGCCTTTTT TAAAAATATA	204
25	TTTGACAGAA ACCAGGTGCC TTCAGAGGCT CTCTGATTTA AATAAACCTT TCTTGTTTTT	210
23	TTCTCCATGG AAAAAAAAA AAAAAAA	212
30	(2) INFORMATION FOR SEQ ID NO: 158:	
	(i) SEQUENCE CHARACTERISTICS:	
35	(A) LENGTH: 1625 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double	

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 158:

CAAAAGATCT ATAATCAGGA CATTGTTTAT GTAAGTTGGA CAANAAAAAT TCTTCCCCTT 60 TATGTCCACC CTTCCTATGA TTGCAAGACA AAATTTCCCT CCTTTACCTC ATCCCTATAA 120 CATGGGAGGC TGAGAAAAAT GAGGGGAGAT GGAACCAGAT ACAAGGAGAT CCAATAAGAG 180 AAGCTTATTT AAATATTGTG AAATAAAGGA AGAMCCAAAG CATTTTTTTA AGTGGGGAAT 240 CCTTTGAAC AGTTATTATT TATCCATATT ATTAAYAACA TCTTTTCTGA CAAAATCCAT 300 CAGATGAAGT GTAAATGGAT AATCTTTTAA TGGATCTAAA CCTAGAAAGT TTCACTTACT 360 GTTCATGTCC GTGTTCCAGA ATTGTGAAAT GGTGTGGGT TTTGCTTTCC AAGTTCTTCT 420 55 CTGCCTCCTC TTAATTCTCT AATTCCATGT CTTACAGAAG AATGAGAAAT TTCTTTCTTA 480 CTTGAGTATC ATGCTCTAAA AAACTTGGCT TCAGTCACAG AAACGCTGGC TCTCCTGTGC TTATATTGAA GCCAACTGCC TTTAATTCTT GGGCCCTCTT ATATTTTTAA GGTGCAAAAT 600

410

	TTGAAGTCTC	AGTCACCAGA	CACAGGTTCT	ATACAATTAA	TGATGAGCTG	GAGAAGTAAT	660
				TACTTTCCGG			720
-							
5	ATTGCCACAT	CTTGCCAGAA	TCCCATCTGA	CACCTTAACT	TIGTCAGGTT	TCCTACAACT	780
	TGCTAATCAA	GTTTTATACA	TTCTAAATCT	CCCCAGTTTC	TTTGGGGCTG	GAAGATGCAA	840
10	CITCCATITA	ATAGAAACTT	TGAAATCTTG	GGGTAAGGGA	GCAGTGGGGG	GACTAGGGAG	900
10	AAGGATAAGA	AATAGAATTA	TTGAAAAGCC	CCCACCAGGG	ACCTTCCTGG	CCAGAATATG	960
	CAGAGTAATT	CCTGCTGGCT	TCACCTTTGA	AAGTCCCTCG	AAACTATGCA	GATGAAACTG	1020
15	AGTCTGTTTT	TGATATTGTC	AGATGTATTC	TACCTTGGAA	GTCCCNACAC	CTAAACTGGA	1080
	ATTCTTGTAT	TTACATCTCC	TCCACTGTCC	CCCACACCAC	CCCTCAATTC	CTGCTGCCCC	1140
20	TGCTAATGTT	AAGCATTTTT	CTCTTGTTAT	CATCAGGTTC	ACATTAAAAM	CAGRTACTTA	1200
20	CAAACTGACT	TGAAGCACAG	ATACTTTTAC	GAATGTGATA	AAATATTTTC	TTAAGAAAAG	1260
	GAAAGAGGAT	GTGGGTCAAA	TAAAACACCG	CATGGATGTT	GATTGGTGAA	TACTGGTGTA	1320
25	AGAAAAGGGA	GCTCAGGAAT	TTTTATTACT	GTATTTGTAA	ATGAGTTTGA	AGGAATTTGT	1380
	AAATGCCACT	GGTACATTTT	TAAGGTGACA	CATTIGCTCC	TTATAAAGTT	TTAAAAATTA	1440
30	ACAGGGTAAG	CTTAAATGAC	GITTGCCAGT	AGTTTTACTT	тататаатса	ATATTGATAT	1500
30	TGTTGCTGAA	CTATGTAACT	TTATGATGCA	TTTTTCAGTC	CCTTTTCAGA	GCAAATGCTT	1560
	TTGCAATGGT	AGTAATGTTT	AGTTTAAATT	GACTTAATAA	ATTMITACCT	GAGCAAAAA	1620
35	AAAAA						1625

40 (2) INFORMATION FOR SEQ ID NO: 159:

45

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1687 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 159:

50 CGGGGTCACC AGTTATTAGA GGAAGTAACA CAAGGGGATA TGAGTGCAGC AGACACATTT 60
CTGTCCGATC TGCCAAGGGA TGATATCTAT GTGTCAGATG TTGAGGACGA CGGTGATGAC 120
ACATCTCTGG ATAGTGACCT GGATCCAGAG GAGCTGGCAG GAGTCAGGGG ACATCAGGGT 180
CTAAGGGACC AAAAGCGTAT GCGACTTACT GAAGTGCAAG ATGATAAAGA ĠGAGGAGGAG 240
GAGGAGAATC CACTGCTGGT ACCACTGGAG GAAAAGGCAG TACTGCAGGA AGAACAAGCC 300
AACCTGTGGT TCTCAAAGGG CACCTTTGCT GGGNATCGAG GACGATGCCG ATGAAGGCCC 360

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	TOGAGATCAG TCAGGCCCAG CTGTTATTTG AGAACCGGYG GAAGGGACGG CAGCAGCAGC	420
_	AGAAGCAGCA GCTGCCACAG ACACCCCCTT CCTGTTTGAA GACTGAGATA ATGTCTCCCC	480
5	TGTACCAAGA TGAAGCCCCT AAGGNAACAG AGCTTCTTC GGGGACAGAA GCTGCCACTG	540
	GCCTTGAAGG GGAAGAAAAG GATGGCATCT CAGACAGTGA TAGCAGTACT AGCAKTGAGG	500
10	AAGAAGAGAG CTGGGAACCC TCCGTGGTAA GAAGCGAASC GTGGGCCTAA AGTCAGATGA	560
	TGACGGGTTT GAGATAGTGC CTATTGAGGA CCCAGCGAAA CATCGGATAC TGGACCCCGA	720
15	AGGCCTTGCT CTAGGTGCTG TTATTGCCTC TTCCAAAAAG GCCAAGAGAG ACCTCATAGA	780
13	TAACTCCTTC AACCGGTACA CATTTAATGA GGATGAGGGG GAGCTTCCGG AGTGGTTTGT	340
	GCAAGAGGAA AAGCAGCACC GGATACGACA GTTGCCTGTT GGTAAGAAGG AGGTGGAGCA	900
20	TTACCGGAAA CGCTGGCGGG AAATCAATGC ACGTCCCATC AAGAAGGTGG CTGAGGCTAA	960
	GCCTAGAAAG AAAAGGAGGA TGCTGAAGAG GCTGGAGCAG ACCAGGAAGA AGGCAGAAGC	1920
25	CGTGGTGAAC ACAGTGGACA TCTNCAGAAC GAGAGAAAGT GGCACAGCTG CGAAGTCTCT	1080
	ACAAGAAGGC TGGGCTTGGC AAGGAGAAAC GCCATGTCAC CTACGTTGTA GCCAAAAAAG	1140
	CTGTGGGCCG CAAAGTGCGC CGGCCAGCTG GAGTCAGAGG TCATTTCAAG GTGGTGGACT	1200
30	CAAGGATGAA GAAGGACCAA AGAGCACAGC AACGTAAGGA ACAAAAGAAA AAACACAAAAC	1260
	GGAAGTAAGC AGAGCTGCCA GGCTCCCAGG AGAGCATGGG GACTAGGAGG AAGGGTGTGG	1320
35	CATGGCTCAG TCTGGCCCCC TTGATTACCG GCCTAGCCCC TGCTCACATC ACAGCTGTCT	1380
	GAAGAACAGT GAGGTGGAGT GCCTAGAACT CCCGTGGTGG TCCTGAGCAG AGAGGAGGAT	1440
	GTCCTCCTGC CTGCCTGAAG GTCTCCCATG AAAACACTGC TGAACTGTGT TGACACTCAT	1500
40	GACCCTTTTT TTAAACCGTT AAAGGGAAGT TCGGTGTTGG AGCGATACTC AATGTAGTCA	1560
	GICTACACCT GGACGIGIGG GCCACTTAAG CCCTCCCCAC CCCCATCCTA TTCCTRAATA	1520
45	AAACCAGGAT AATGGAARAA AAAAAAAAAA AAAAAAAAAG GGGGGGCCCN TAAAGGGNCC	1580
73	CANNITT	1687

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(2) INFORMATION FOR SEQ ID NO: 160:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1842 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 160:

60

	GGATGACAGA	TTGCGACANA	GATTTGTGAC	CCTTCCTGCT	GAACTTCAGA	GGGAGCTGAA	60
	ANCAGCGTAT	GATCAAAGAC	AAAGGCAGGG	CGAGAACAGC	ACTCACCAGC	AGTCAGCCAG	120
5	CGCATCTGTG	CCCCGAGAAT	CCTTTACTTC	ATCTAAAGGC	AGCAGTGAAA	GAAAAGAAAA	180
	GAAACAAGAA	GAAAAAAACC	ATTOGTTCAC	CAAAAAGGAT	TCAGAGTCCT	TTGAATAACA	240
10	AGCTGCTTAA	CAGTCCTGCA	AAAACTCTGC	CAGGGGCCTG	TGGCAGTCCC	CAGAAGTTAA	. 300
10	TTGATGGGTT	TCTAAAACAT	GAAGGACCTC	CTGCAGAGAA	ACCCCTGGAA	GAACTCTCTG	360
	CTTCTACTTC	AGGTGTGCCA	GCCTTTCTA	GTTTGCAGTC	TGACCCAGCT	GGCTGTGTGA	420
15	GACCTCCAGC	ACCCAATCTA	GCTGGAGCTG	TTGAATTCAA	TGATGTGAAG	ACCTTGCTCA	480
	GAGAATGGAT	AACTACAATT	TCAGATCCAA	TGGAAGAAGA	CATTCTCCAA	GTTGTGAAAT	540
20	ACTGTACTGA	TCTAATAGAA	GAAAAAGATT	TGGAAAAACT	GGATCTAGTT	ATAAAATACA	600
20	TGAAAAGGCT	GATGCAGCAA	TCGGTGGAAT	CGGTTTGGAA	TATGGCATIT	GACTITATTC	660
	TTGACAATGT	CCAGGTGGTT	TTACAACAAA	CTTATGGAAG	CACATTAAAA	GTTACATAAA	720
25	TATTACCAGA	GAGCCTGATG	CTCTCTGATA	GCTGTGCCAT	AAGTGCTTGT	GAGGTATTTG	780
	CAAAGTGCAT	GATAGTAATG	CTCGGAGTTT	TTATAATTTT	AAATTTCTTT	TAAAGCAAGT	840
30	GTTTTGTACA	TTTCTTTTCA	AAAAGTGCCA	AATTTGTCAG	TATTGCATGT	AAATAATTGT	900
	GTTAATTATT	TTACTGTAGC	ATAGATTCTA	TTTACAAAAT	GTTTGTTTAT	AAAGTTTTAT	960
	GGATTTTTAC	AGTGAAGTGT	TTACAGTTGT	TTAATAAAGA	ACTGTATGTA	TATTTGGTAC	1020
35	RGGCTCCTTT	TKGTGAAYCC	TTAAAAACTC	AACTCTAGGA	RGCAACTACT	GTTTATTATA	1080
	CTAAARGGCT	GAAAAMCCTC	CYCÉCCYCYC	TGCTAAGCTC	TGAAATYCCT	GAGAGGTCTC	1140
40	AGACCGGGAT	TCTACTTGTT	CCAAGAAAGG	GTAAAGCTTC	TAAACCATCT	TATTCTTGTC	1200
	TCCAAGCATG	AACACAGGAG	CATGTYAAGA	AAATCTTTAC	TACTTTCTYC	CATGCGGAGA	1260
	AATCTACATA	TTTTGAATTA	GAAACACCCT	CACACCCACT	TGAAGATTTT	TTTCCTGGGA	1320
45	ACATTATGTC	CCGTAGATCA	GAGGTGGTGT	TGTCTTTTTG	CTTCTACTGG	CCATTGAGAA	1380
	ACTITGATGA	TAAAAAAGAA	COGTATAGAT	TTTTCAAACG	TATATAAAAT	ATTTTTATGT	1440
50	TATATGTTAT	GCCATAACTT	TAAAATAAAA	ATAGTTTAAA	ATTCTATGCT	AGTGGATATT	1500
	TGGAACTTTT	TCCTCAAACA	AACACCCCAC	ACTGACTTCA	GCAAAACCCT	AAAACTAGCT	1560
	ACAGATTACT	ACTACGAATG	AATCATYAAG	TTTTGTGTCT	GCAACAATTT	AGAAGCACTA	1620
55	AGCCCAAATA	TCAGGAAATG	TGTGTATGAT	GGAATTTTCT	AGGACAAAAC	AGATCAAGAT	1680
	TAAAACAGGA	TCAAGGATTA	ATGGTATAAA	AATGGTCTAC	TAAAACAGGA	TCAAGGATTA	1740
60	AAACAGGATO	AAGGATTAAT	GGTATAAAAA	TCTCTACTGG	TTACCGGGTG	GCNGGGCCAT	1800
00							

	ACACGGTAGT CGTCGATCGA TAGTTTAGTT TCGNAAGGGT AA	1842
5	(2) INFORMATION FOR SEQ ID NO: 161:	
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 770 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	-
15	(xi) SEQUENCE DESCRIPTION: SÉQ ID NO: 161:	-
	GOCACGAGCC CTATGCTGTT CTTGTGATAA TGAGTGAGTC TCACAAGATC TGGTGGTGTT	60
	ATAGGCATCT GGCATTTCCC CTGCTGACGC TCATTCTCTA TCCTGCCACC CTGGGAAGAA	120
20	GRGTCTTCTG TCATGATTGT AAGTTTCCTG AGGCCTCCCC AGCTATGTAG AACTGTGAGC	180
	CAATTAAACC TCTTTTCTCT ATAAATTATC CAGTCTTATA TATTTCTTCA TAGCAGTGTG	240
25	AGAACAGATA ATACCGTAAA TTGGTATCAC AGAGAGTGGG GTGTTGCTAT AAACACATCT	300
25	GAAAATGTTA AAGCAAATTT GGAACTGGGT AACAGGCAAA GGCTGGAACA GTTKGAAGAA	360
	CAGTTAAGAA GAAGACAGGA AAATATGAGA AATCTTGAAA CTTCCTAGAG TCTTAAAGGT	420
30	CTCAGAAGAC ATGAAGATGT GGGAAGCTTT GGAACTTCCT AGAGACTTGT TTGAATGGCT	480
	TTGACCAAAA TGCTGATAGT GATATGGACA ATGAAGTCCA GGCTGAGCTT ATCCAGACAG	540
	ACATAAGAAG CTCGCTGGGA ACTTGAGTAA AGATCACTCT TGCTAGGCAA AGAGACTGGT	600
35	GGCCTTTTTT CCTCTGCCCT AGAGATCTGT GGAAATCTGA ACCTGAGAGA GATGATTTAG	660
	GGTATCTGGC AGAAGAAATA TCTAAGCGGC AAAACCTTCM AGAGGAAGCA GAGCATAAAC	720
40	GTTTGAAAAA TTTGCAGCCT GACNATGGGA GACCAAAGTT AAACCCAATT	770
45	(2) INFORMATION FOR SEQ ID NO: 162:	
	(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 519 base pairs(B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 162:	
55		60
55		120
	TGGGGTCACA GACTTCARAC CTGATGACCT GGGCTCAGAT CCCAGCTCTG CACCTACCAG	
	COGTIGTGACA AGGTGTCCTC TCTGAGCCTC AGTCACACAC TGCCTTAACG GTTGGGCCTC	18

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	ATGGAGCTGT TTGTGAAGGT TAAATGGGAA GACATAAAGC ACTTAGCCCA GAGCCAAGGA	240
	CATECTGAAT AGGATAATGG TGGCCTCCTT TGGCGCTGTG CTGGTGCAGG TGTGCCGAGG	300
5	AAYTGGGCAG GGGTGACAGA TACCTCTTCT AACCTAGTTC CTTTCCAAGA ACCTAATTGG	360
	TGTCTCTCCC TCCCCCAGGC AATTGGAAGG AGGAGGCTGG GCCCCAGCCC CAGAATACGG	420
	GAGGTTTCTC ACCGTGGTAG GGAAATTGCT GGGTTGGGGG TGTGGGCAAC CACAGTGATC	· 480
10	GTCTCTCTGC AGGACGGATG AGGCTTTGCT GACAGAGGC	519
15	(a)	
	(2) INFORMATION FOR SEQ ID NO: 163:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 753 base pairs	
20	(B) TYPE: nucleic acid	
	(C) STRANDELNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 163:	
25	GGCACGAGCG GCACGAGCAG CCAGTTGCTG ACTGGCACAT GGCCTCCAGC GTCCCGGCTG	60
	GTGGGCACAC TAGAGCCGGA GGGATCTTCT TAATTGGTAA ATTGGATCTT GAAGCTTCAC	120
30	TGTTTAAATC TTTTCAGTGG CTTCCCTTTG TACTTAGAAA AAAATGCAAC TTCTTCTGCT	180
	GGGACTCATC CGCTCACAGC CTTCCCCTCC ACCCTCTCTC TGCCTCATGC TCTGCCCCTG	240
25	CCTGCCATGC CTCCGATACT CACCTTTTGT ACCCCAGCAC CCGTGCCCTC TGCCCCTCGA	300
35	TCTTTGCCTG GCTGGTTGCT CCTCACTCAG TGTTCAGGAC AAATGCTCCT GGCCCTACCC	360
	CATCTAGCCA GTCTAGCCCG GTCTTCCCTG TCTTCCCTGT TTCATTCATG GCTCTTATTG	42
40	TTTGTTWACT TGTGTGCTGT TGACTTTTAA CTCTCTCAGT CCCCACTGGA ATGCAAGCGA	48
٠.	TCTCCCAAGC TCCTAGAATT GTTCCTGCCT CTTCACAGGC CCTTACGCTG TGTGTGCTCG	54
45	TGCCGAATTC GGCACGAGGG TATGTGCACT TGCTGGTATG TATGTAGGTG TTTGCTAACA	60
43	CATACGTGCA CACGCAGAAT GCTTCCAGGG GACTGCACAG CCTCTAGTTC GCAGCCCCCA	· 66
	CCCCTCCCTT TGSCCCTGCA CTCTCCCCTC TCTGAGCTGC ATTCGCATGA AAGGGTGCAN	72
50	GGTTCCTGAN CCCGCNAGCG NCACCTCCTG GGA	75
55	(2) TITOTHORNEY POD 670 TD NO. 164	
JJ	(2) INFORMATION FOR SEQ ID NO: 164:	

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1400 base pairs

(B) TYPE: nucleic acid

60 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 164:

5	GGCACAGTTT ATTANTACCT ATTATGGGAA AGTCACTTTG GTTGGCATTG AAAATTACAT	60
	CATCTITAAA GCAGTATTIG TCCCCAGATG GACTCATCAC TAGCAAAGAC TAGGITCATT	120
10	GGAAGGCATA GGGTGAGAGA ATGGGAAGAT GRAGTGGAGG CGGGTTGTTA AAGTGCTGTC	- 180
10	AGTGAGTGAT TTTGTCTACT TGAACAATGG TCCATGTTTG GGGGCATATT GTGTTTCATA	240
	AGAAGTGAAA GGTATTTGCA AAGTAAGCTA CAAATGACCC ATAAATCTGT TAACAACAGT	300
15	CCTTAATATG CAAAGATGAA AAACAAGCAT TACTGCTACC CAAAGGGAAC TGGTGCTTGG	360
	TGATGTGCAG ATGGGGCTGT TGGTTAAGAG AGCTATTACA GGTTTTCTCT CTTAGGTTTC	420
20	ATAGGAGGTA GTTACTGAGA TGAGATTGTT TTATCTTTTT GAATACAGAT CTCTTGTCTT	480
20	GAGTTAGTTC TGAGGATGGG AGTAATAAAG GAGTTTTTTG TTTTTTTGTT TGTTTGTT	540
•	TTTTGGCTCC TTAGTAATAC TCCTCTGACA TTTATTTCTA TTATTCTTCA AAGAAAGGAA	600
25	ACCAACTGAA ATGTTTGCTT TAACAAACAT TTTAATAAGT TCTCTGGGTT TTTTTTTCCC	660
	CTTTTAAAAA AATTAGCATA TACCATAGCA ATAAAAGAAC TAATGTTAAC TATTGTATGC	720
30	TACAACTTAA GTGATTTTC TAAAGAAGCA CAATGTCATT GRAAGTATTA TTGAAAAGGA	780
50	TCATAGTCAC ATTGAATTTG TGAAGGCCAA AGAAATTGAA GGGAGTGATA TTTTCATTTT	840
	ATGATATTCA CATATTTAGT AAATTTTGTG TACAAGAATA CCAGGCAGAG TGTTTTACCC	900
35	ATGGAAACAG GTTTCAGATT ACTTTGTTTT TACTGTTAGA GTCTCAAGTT TAGAAATGCT	960
	AACACTTAAA TCAGTTTTT TCTÇACTATA CTTGAAGATT GTTAATATTT TGATATCTTC	1020
40	CTAGCTTGAT GGAATTTAAA CATATCTTCA GATCTGTGAC AGTGACAGCC AATAGGACTG	1080
40	ATAATATTAG CTTCAAACCA ATAATATCCA GGGTTAAAAT AAAAATCATA GTGAAAGTAC	1140
- ,	GATTGTAAAA TTATGCTATA TTAACTTTTA AGTCTGTAAT AACTTGACAT CAAAATGTTA	1200
45	TGTAATTACC ATAAATAATG GCTAGCGAGA ACATCTTTGG AAATTCTCAA ATTACCTTTC	1260
	TTACTACACT GTTTGCAGAA TGAATGTAGA AATGATCCTG TTAGCTTTCT GAATGTTCTG	1320
50	TGGTTGAATG TGTTTTGCT TAAATAAAGC TTTTGGTATT TGTTTAAATW ACAAAAAAAA	1380
50	AAAAAAAAA AAAAACTCGA	1400

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(2) INFORMATION FOR SEQ ID NO: 165:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2153 base pairs
(B) TYPE: nucleic acid

416

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 165:

)	CAGGCCTCAG GOCCTCTGGT GGCTCTGGCC CAGACAGTAT TTGCAGTTCT TGTGCTATGG	60
	GTGGGAGTCT TCTTCCTCAA GTTTCGGCAG CTGTGCTGTG	. 120
10	CAGGGCTCAA GGGCTGTGGT CCGCTCAGGG TCTCATTTCC CCAGGCCAAG TTCAAGGCAG	180
	CAGCCCTTTG TGAGGCGCTC TTGGCCCTGG GCTGGAGGGA GAACTTTAAG CTTTTTTGCT	240
15	CACAGGGACG TGGTATGGGC CCTGGGTGCA GGTGCCCACA TTCTGCTAAT GAGAGCTTTG	300
13	TCTGATCAGT CCTGGGTCCA TCAGTTTGTC CATGTGTCCG GCTGCCAGCC CGTCCCTTGG	360
	GATCCTTCCC CTGGGGTGTA GCCTTGTTCA TTAGTATATA CTCATTCCTT CATGCTTTCC	420
20	TCAGCAGAAC ACTTCCACTT CTGAGGTGAG CTTTTGCCCC RTGCCCTTCC TCCACAGGTG	480
	TTGCCTTTTT ATAAAGACCT GATAGCAGAA TAAATTGGTG TTTCCCTGTT GACCCAGCAC	540
25	CATTICTGTG GGCCTAGAAT ATGGCCCTCA ACCCTTAGAG TGGGGCAGTG AGGGCTTGAG	600
25	GAGTGACCCT TCCTTTCTCA TGGTTTTAGT CATTTTGGCT GCCAGCCCTT AATGGCACAG	660
	ATCTGCTGCT TCTAACAGAT GGCCAGGAGG TGACACCGAT TTCAGCCATT GCCAAGGTTA	720
30	GCACCCTCTC CTTTGAGCCT AGGGCCACAC TGTTCATTGT CACTTTAGGC AAGTGCCTGT	780
	TTGGCTTTAA AGGTAAGCCT GCCAGCTGTG AGAAGCCTTG GTAACTGATG GACTCATTTC	840
	CTGGTCCTTA AAGATGCAGC CTCTTAAGGG CTCCTTGATG GATGCCATCT CTCCTAGCCC	900
35	CCAGCCCTGG TGCCACTGGT GGGCAGGTTC CCATTCTTTG GGGCTGGGAG GGACAGCTTG	960
	CCTGTTTCTG GTCACAAATT ACAGTCTTCT CTCCTGTACC ATTCTGTGGC TTCAGCATGG	1020
40	GGGCAGTAGC CTTTCATTAG TGTAGATAGT CATTCCCTGG TAGGGTGGAG GGTAAGACAT	1080
	AGGGTCTGGA ACTGTTTGGG ACCTTTTGGG GATGTCCTGT GCCTCCCAGA TTCCTMGATT	1140
	CTGGGAGGAG AGGCTGCCGC ATTCTGCTGC TCCTCACAGC GAGCAAAGCT GCACCCACTT	1200
45	ACATTCAGTA TTTTCCTGGC ACTACAAAGA GTGGGAAGGC CTGGGATTTG CTGCTGCTCC	1260
	CTTAGAGCAG GGCCCCTYTT TTCAGCACTT TGGACACCTG GAGACCCAGC CCTGTTATTT	1320
50	AATGGTAGTG GGCAAGTGTG TGTGCATACT GTCTGCCACT GCTTTCTCCC TGCCCCATGC	1380
	CAGAGAGCCC TGTCCCTGCC AGGCCCAGCC TTCTTAGCCC CAACTTGGGA ACAAAGTGCA	1440
	ACATGGGATC ATGGGTTGGG GTGCTCAGGT GAGCCCTCTC TATAGTGCTT CCCTGGGCCA	1500
55	AGCTGACACC AGCCCCTGAG GGTGGGGTGG GACGGGTGGT GCTTAAAAGA GGAAGGGGAC	1560
	CAGTGTAGCA ACTTGCCAGG GACCCCACCC CTCCCTCTCT GGGCCTGTGC AGTGAGCATG	1620
۲0		1680
60	GOGATTOCCA TCAAGGGGCC TGGCACCTGT GCTAGTTACG TAGCCGCTGN TCACGCGCTC	1090

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	ACTCCTGACC	ACATGCACGT	TOCCTAGATG	CAGACTGCTT	TGAACTTTAA	AGCTGTACAA	1740
5	TTTGGTTATG	TTTGTGCTGA	CTTAAAATAT	attitaatga	GGAAAAAATA	ATGGAGAACC	1800
3	CTGGGAAGGA	CCTGGTTCTT	TIGCTICICG	GGGAACTGTA	AGCCCTCGCG	TTCTGGGAAT	1860
	CGCTCTCTGC	TGCTCTTTCC	TGGAAGCTAA	OCCIGICICC	ACCGCCCGAG	GCCTGCGGCCG	1920
10	GTGCTCCCGC	CGCAGTTGCG	TTTGCTTTGG	ACCTTGCGTG	CGGGGGAGGG	GGTGCTCGGT	1980
	CCGAGCCCGC	TCCTTTCTGT	ACACCTAGCG	CICCCCCCC	CCCTTCTCTC	TGAGGTCGTG	2040
15	TATGTCAAAA	ATAAAGCCGC	TAGAAACGGA	АААААААА	ааааааааа	AAAAAAAA	2100
15	AAACTCGAGG	GGGGGCCCGT	ACCCAATTAA	CCCNNTATGA	TCTATAAAGC	GTC	2153

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(2) INFORMATION FOR SEQ ID NO: 166:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1251 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 166:

30 GCCCACGCGT CCGCCCACGC GTCCGGCGGT GCGGAGTATG GGGCGCTGAT GGCCATGGAG 60 GECTACTECC CCTTCCTGCC GCTCCTGGGG TCGGCACTGC TCGTCGGCTT CCTGTCGGTG 120 ATCTTCGCCC TCGTCTGGGT CCTCCACTAC CGAGAGGGGC TTGGCTGGGA TGGGAGCGCA 180 35 CTAGAGTTTA ACTOGCACCC AGTGCTCATG GTCACCGGCT TCGTCTTCAT CCAGGGCATC 240 GCCATCATCG TCTACAGACT GCCGTGGACC TGGAAATGCA GCAAGCTCCT GATGAAATCC 300 40 ATCCATGCAG GGTTAAATGC AGTTGCTGCC ATTCTTGCAA TTATCTCTGT GGTGGCCGTG TTTGAGAACC ACAATGTTAA CAATATAGCC AATATGTACA GTCTGCACAG CTGGGTTGGA 420 CTGATAGCTG TCATATGCTA TTTGTTACAG CTTCTTTCAG GTTTTTCAGT CTTTCTGCTT 45 480 CCATGGGCTC CGCTTTCTCT CCGAGCATTT CTCATGCCCA TACATGTTTA TTCTGGAATT 540 GTCATCTTTG GAACAGTGAT TGCAACAGCA CTTATGGGAT TGACAGAGAA ACTGATTTTT 600 50 TCCCTGAGAG ATCCTGCATA CAGTACATTC CCGCCAGAAG GTGTTTTCGT AAATACGCTT 660 GOCCTTCTGA TCCTGGTGTT CGGGGCCCTC ATTTTTTGGA TAGTCACCAG ACCGCAATGG 720 AAACGTCCTA AGGAGCCAAA TTCTACCATT CTTCATCCAA ATGGAGGCAC TGAACAGGGA 780 55 GCAAGAGGTT CCATGCCAGC CTACTCTGGC AACAACATGG ACAAATCAGA TTCAGAGTTA 840 AACAGTGAAG TAGCAGCAAG GAAAAGAAAC TTAGCTCTGG ATGAGGCTGG GCAGAGATCT 900

•	ACCATGTAAA	ATGTTGTAGA	GATAGAGCCA	TATAACGTCA	CGTTTCAAAA	CTACCTCTAC	960
	AGTTTTGCTT	CTCCTATTAG	CCATATGATA	ATTGGGCTAT	GTAGTATCAA	TATTTACTTT	1020
5	AATCACAAAG	GATOGTTTCT	TGAAATAATT	TGTATTGATT	GAGGCCTATG	AACTGACCTG	1080
	AATTGGAAAG	GATGTGATTA	ATATAAATAA	TAGCAGATAT	AAATTGTGGT	TATGTTACCT	1140
10	TTATCTTGTT	GAGGACCACA	ACATTAGCAC	GGTGCCTTGT	GCAKAATAGA	TACTCAATAT	· 1200
	CTGAATATCT	GTCTACTAGT	AGTTAATTGG	ATAAACTGGC	AGCATCCCTG	A	1251

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(2) INFORMATION FOR SEQ ID NO: 167:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 882 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 167:

GACSMTCTAG	AACTATGGTC	CCCCGGGACT	GCAGGAATTC	GCCACAGCGG	CICCGGGGGG	60
GAGGTGAGGG	GCGCGAGGTT	CCCAGCAGGA	TGCCCCGGCT	CTGCAGGAAG	CTGAAGTGAG	120
AGGCCCGGAG	AGGGCCCAGC	cccccccc	CAGGATGACC	AAGGCCCGGC	TGTTCCGGCT	180
GTGGCTGGTG	CTGGGGTCGG	TGTTCATGAT	CCTGCTGATC	ATCGTGTACT	GGGACAGCGC	240
AGGCGCCGCG	CACTICTACT	TGCACACGTC	CTTCTCTAGG	CCGCACACGG	GCCCCCCCCT	300
GCCCACGCCC	GGGCCGGACA	GGGACAGGGA	GCTCACGGCC	GAYTCCGATG	TCGACGAKTT	360
TCTGGACAAK	TTTCTCAGTG	CIGCCGIGAA	GCAGAGTGAC	YTTCCCAGAA	AGGAGACGGA	420
GCAGCCGCCT	GCGCCGGGGA	GCATGGAGGA	GAGCGTGAGA	RGCTACGACT	GGTCCCCGCG	480
CGAMGCCCGG	CGCACCCAGA	CCAGGGCCGG	CAGCARGCGG	ANCGGAGGAR	CGTGCTGCGG	540
GGCTTCTGCG	CCAAYTCCAG	CCTGGCCTTC	CCCACCAAGG	AGCGCGCATT	CRACGACATC	600
CCCAACTCGG	AGCTGAGCCA	CCTGATCGTG	GACGACCGGC	ACGGGGCCAT	CTACTGCTAC	660
GTGCCCAAGG	TOGCCTGCAC	CAACTGGAAG	CGCGTRATGA	TCGTGCTGAG	CGGAAGCTGT	720
GCACCGCGTG	CGCCTACCGC	GACCCGYTGC	CINTCCCCCCCC	GAGCACGTGC	ACAACGCCAG	780
CGCGCACTGA	CTTCAACAAT	TCTGGCGCCG	CTACGGGAAG	TCTCCCCCAC	CTCATGAAGT	840
CAAGCTCAAG	AATACACCAA	TTCTTTCTGC	GCGACCCTTC	TG		882

⁽²⁾ INFORMATION FOR SEQ ID NO: 168:

419

(i)	SEQUENCE	CHARACTERISTICS:
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(A) LENGTH: 1208 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

5 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 168:

10	GGGAAACTCA	AAAGGATGAT	GGAATGGTTG	ATGGAGCCAG	AGCCTAGAAG	TRAAGGGATA	. 60
10	CAGAGTGAAG	ATAGAGGTAT	TTACGTATAT	ATTATAAWIT	GCTTTGGAAT	TACGTAGGGA	120
	TTCTTAAGAA	AAGATCATGA	CAGGACAGCC	ACATTTGGTA	AAATGTCAGG	GCAGCCAGTG	180
15	CATGGTCCTC	CIGGGGCTCC	TCAGTTGACG	GGTTTAAATC	ATTTCCTGAT	CCCCCTGCCC	240
	TGGTTTGAGG	AATGCATACA	GTACGTGAAA	TGCCTGTGGT	ATGAGTTGCA	ATGGGCAATC	300
20	AACCTGGGTA	AATCCAAGAT	TAATGATTAG	TTCTAAAGAT	CCAGTTGAAG	TTCTAGAGTG	360
20	GGAATTTTCC	GTCAAGCARC	TCAGCACAGC	TTTATGCCTG	TTCCTCTAAT	AACGATAGGT	420
	AACAAATAGC	TGTGTKTWCA	CAGCTAGGAR	GATAACCAAA	TCTAGAGTTC	TTGARTCTCA	480
25	TTTAATAAAT	aaktattatg	AGTACCAACT	GCATATTTCA	GGCACTGCAT	TIGACTCTGT	540
	TAAATACTGA	TYCCTTAKGA	CMSCCACWIC	AGAWAACMIT	AATCTGTCTG	ATCAATAAAC	600
30	AGCTTGACTT	AGAGRGGTAA	AATAGCTTGC	CACAGGTWAC	CCAATTAGTA	GGTAACAGCG	660
30	ACAGAATAAC	agtgcagtta	AAATCTTAGA	CTGGAGACTA	ATTGCATAAG	TTTGAATTTC	720
	AGTTCTGCTA	TGTAAATTTG	GGTGAGTACC	TTAATTYACC	TGAGTCTCGG	TCTTTATATC	780
35	TGTAGAATGG	AGCTAATGAT	ATTACTTAAT	TTGCTTTATG	TGAGATTAAA	TGTACTAATA	.840
	TATGTAAATC	ACTTACAACA	GCAŢŢŢĠĀĊĀ	TATTTGACAT	ACTTAATATA	TTTGCTACTA	900
40	ATACTATTAG	CAACAGCATT	CTGATTTTCC	AAGTTGAAAT	TCAGTGTTTT	CTTTTTTACT	960
40	TTGCCATAAT	TTACAATGTT	GTGCTCTGTA	AACCATAAAT	TTCCCTGAGG	TGTTGTCAGG	1020
-	ТТААААААА	ATCACTATGG	CCCCCARNIMA	CTTGGAAAAT	AGAAATGAGA	CCAGCTTCAT	1080
45	CTATATTCTT	TACTGCAAAT	AACTTAGAAT	TGTAATAGGC	TAATATGTAC	TGGGACTTCC	1140
	AATTTGGGAA	TATGACAAAA	ATAATACTAT	TTAGCTAAAA	CATATACAGA	ACTTATTTTT	1200
50	CCTCTGAA						1208

(2) INFORMATION FOR SEQ ID NO: 169:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1307 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi)	SPOUENCE	DESCRIPTION:	SEO.	TO NO.	169
\^_/	SECULIACE	DESCRIPTION:	SEL	m wo:	103

5	GGCACGAGAG	AAAAGAGGTT	GAGAATGTTT	TCTAGCAGGC	AGAATGTGCA	TACATGTTTT	60
J	CATGARTGTC	CTTTGGGTGC	TGTTTCTTTT	AAATCCTCTG	TGCACAGGGC	TCTGGCCTTT	120
	ARTAAACTGT	TTTTCTGTCT	TACGTCATGC	TGACTGGGTG	CTAGGGGCTG	ATTACAAAGG	180
10	GGAAGAGTTG	AACAGACATC	AGGGGCCGAT	GAAACCAAAG	GACTAGGAGT	CAGGAGAACA	240
	AGTCAGGGAT	TAGGAGACAG	CGGTTTGGTT	TATTGTTATC	CAGCTGGAGG	ACTCCTAGGG	300
15	GCAGCAGCAG	GAGGAATACC	AGGCCACGG	AGGGGCAGGA	GTCTCACAGT	GGAGGGCAGA	360
13	CTCTAACAGA	TGCCAGCTGA	ACCCTCCCTG	GCCCTGGATG	TCATACGAGT	TGGGGACCAG	420
	AAATCTGGGC	TCAGAGAACC	CGTCCAGGGA	GATTTGAAGC	CATGGGTTAT	CTTCTAGAGT	480
20	TGATACTGAT	TTTTATATAA	AATTTTTATT	GATGTTTAAT	ACCTTCTGAA	ACAGGAGGGT	540
	AAGATCAGAT	GGGAAGCCCY	TCTGTTGAAG	GATCTTGGGA	ACCTTGGTGG	TTTTTTTTTT	600
25	TIGGTTTTT	TTTTTTTGAT	CGACCTGTGG	ACATCCTTCT	TAATTCGATT	NTGAGGATTT	660
23	GTTTAACTAA	AAAGTTCCCA	AACACAGAAA	GGGCCTCCCC	ACCTGCTTTG	GGGAGCTGTC	720
	TGTSCTGGGA	GTGCCAGGCA	TCCSATGGGA	CCCATCACTG	CCAGTGTCTG	TGCCTCCCAG	780
30	AGGTCAGCCC	TCTCTCTCCC	CTGGCTCTGT	CTCCTCTGTG	ACAGGGCAGA	GCATTTCTGG	840
	TCAGTTTCTC	CATGGTGCCT	CCCACCCCTT	TGTAAAGTGG	ATGGACATGA	TGGAATTCAG	900
35	TTGTCTCACC	CTGATACCCT	GGGTGTTGAT	ATTCACTTTA	CCCGCACTCA	GACACAGGCG	960
33	ACCTTGAAGC	AGTTCTCGGT	GTGTAGAGTC	CACGTGACAG	TCCCCACAGC	CTCCCCAGAT	1020
	AGCTGTGTGC	CTGTGCGCTA	стестетесс	ATTTTCCCAA	CITNGGCGTT	TCACTAAATG	1080
40	CAGCTGATCT	CTCTCTCTGT	GCACTCGTGA	TCCATGTTGA	ACAATACATG	TAGGITCITT	1140
	TTCCACGCAA	TGTAAGAACA	TGATATACTG	TACGTTGGAA	AGCATTTACC	TTATTTATAT	1200
45	ACCTGAATGT	TCCTACTACA	CAAATAAACA	TATATTAAAT	WCTAAAAAAA	АААААААА	1260
7.7	CTGGAGGGGG	GCCCGGTAC	CCAAATCGCC	GGATAGTGAT	CGTAAAC		1307

(2) INFORMATION FOR SEQ ID NO: 170:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1624 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 170:

60

	GGCACGAGGT	cecceccece	GCCGCCTGGA	ATTGTGGGAG	TIGIGICICC	CACTOGGCTG	60
	CCGGAGGCGA	AGGTCCCTGA	CTATGGCTCC	CCAGAGCCTG	CCTTCATCTA	GGATGGCTCC	120
5	TCTGGGCATG	CTGCTTGGGC	TGCTGATGGC	CCCCTCCTTC	ACCTTCTGCC	TCAGTCATCA	180
	GAACCTGAAG	GAGTTTGCCC	TGACCAACCC	AGAGAAGAGC	AGCACCAAAG	AAACRGAGAG	240
10	AAAAGAAACC	AAAGCCGAGG	AGGAGCTGGA	TGCCGAAGTC	CTGGAGGTGT	TCCACCCGAC	. 300
10	GCATGAGTGG	CAGGCCCTTC	AGCCAGGGCA	GCTCTCCCT	GCAGGATCCC	ACGTACGGCT	360
	GAATCTTCAG	ACTGGGGAAA	GAGAGGCAAA	ACTCCAATAT	GAGGACAAGT	TCCGAAATAA	420
15	TTTGAAAGGC	AAAAGGCTGG	ATATCAACAC	CAACACCTAC	ACATCTCAGG	ATCTCAAGAG	480
	TGCACTGGCA	AAATTCAAGG	AGGGGGCAGA	GATGGAGAGT	TCAAAGGAAG	ACAAGGCAAG	540
20	GCAGGCTGAG	GTAAAGCGGC	TCTTCCGCCC	CATTGAGGAA	CTGAAGAAAG	ACTTTGATGA	600
20	GCTGAATGTT	GTCATTGAGA	CTGACATGCA	GATCATGGTA	CGGCTGATCA	ACAAGTTCAA	660
	TAGTTCCAGC	TCCAGTTTGG	AAGAGAAGAT	TGCTGCGCTC	TTTGATCTTG	AATATTATGT	720
25	CCATCAGATG	GACAATGCGC	AGGACCTGCT	TTCCTTTGGT	GGTCTTCAAG	TGGTGATCAA	780
	TGGGCTGAAC	AGCACAGAGC	CCCTCGTGAA	GGAGTATGCT	GCGITTGTGC	TGGGGGCTGC	840
30	CTTTTCCAGC	AACCCCAAGG	TCCAGGTGGA	GGCCATCGAA	GGGGGAGCCC	TGCAGAAGCT	900
	GCTGGTCATC	CTGGCCACGG	AGCAGCCGCT	CACTGCAAAG	AAGAAGGTCC	TGTTTGCACT	960
	GTGCTCCCTG	CTGCGCCACT	TCCCCTATGC	CCAGCGGCAG	TTCCTGAAGC	TCGGGGGGCT	1020
35	GCAGGTCCTG	AGGACCCTGG	TGCAGGAGAA	OGGCACOGAG	CTCCTCCCCC	TGCGCGTGGT	1080
	CACACTGCTC	TACGACCTGG	TCACGGAGAA	GATGTTCGCC	GAGGAGGAGG	CTGAGCTGAC	1140
40	CCAGGAGATG	TCCCCAGAGA	AGCTGCAGCA	GTATCGCCAG	GTACACCTCC	TGCCAGGCCT	1200
	GTGGGAACAG	GGCTGGTGCG	AGATCACGGC	CCACCTCCTG	GCCTGCCCG	AGCATGATGC	1260
•	CCGTGAGAAG	GTGCTGCAGA	. CACTGGGCGT	CCTCCTGACC	ACCTGCCGGG	ACCGCTACCG	1320
45	TCAGGACCCC	CAGCTCGGCA	GGACACTGGC	CAGCCTGCAG	GCTGAGTACC	AGGTGCTGGC	1380
	CAGCCTGGAG	CTGCAGGATG	GTGAGGACGA	GGCTACTTC	CAGGAGCTGC	TOGGCTCTGT	1440
50	CAACAGCTTG	CTGAAGGAGC	TGAGATGAGG	CCCCACACCA	GGACTGGACT	GGGATGCCGC	1500
	TAGTGAGGCT	GAGGGGTGCC	ACCOTGGGTG	GCTTCTCAG	GCAGGAGGAC	ATCTTGGCAG	1560
	TOCTOSCITO	GCCATTAAAT	GGAAACCTGA	AGGCCAAAAA	. AAAAAAAAA	AAAAAAAAA	1620
55	AAAA						162

^{60 (2)} INFORMATION FOR SEQ ID NO: 171:

422

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 2003 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 171:

10	GGCACGAGCC AGCTTGCAGG AGGAATCGGT GAGGTCCTGT CCTGAGGCTG CTGTCCGGGG	60
	COSGTGGCTG CCCTCAAGGT CCCTTCCCTA GCTGCTGCGG TTGCCATTGC TTCTTGCCTG	120
15	TTCTGGCATC AGGCACCTGG ATTGAGTTGC ACAGCTTTGC TTTATCCGGG CTTGTGTGCA	180
15	GGGCCCGGCT GGGCTCCCCA TCTGCACATC CTGAGGACAG AAAAAGCTGG GTCTTGCTGT	240
	GCCCTCCCAG GCTTAGTGTT CCCTCCCTCA AAGACTGACA GCCATCGTTC TGCACGGGGC	300
20	TITCTGCATG TGACGCCAGC TAAGCATAGT AAGAAGTCCA GCCTAGGAAG GGAAGGATIT	360
	TGGAGGTAGG TGGCTTTGGT GACACACTCA CTTCTTTCTC AGCCTCCAGG ACACTATGGC	420
25	CIGITTIAAG AGACATCITA TITTICTAAA GGIGAATICT CAGATGATAG GIGAACCIGA	480
23	GITGCAGATA TACCAACTIC TGCTTGTATT TCTTAAATGA CAAAGATTAC CTAGCTAAGA	540
	AACTTCCTAG GGAACTAGGG AACCTATGTG TTCCCTCAGT GTGGTTTCCT GAAGCCAGTG	600
30	ATATGGGGT TAGGATAGGA AGAACTTTCT CGGTAATGAT AAGGAGAATC TCTTGTTTCC	660
	TCCCACCTGT GTTGTAAAGA TAAACTGACG ATATACAGGC ACATTATGTA AACATACACA	720
35	CGCAATGAAA CCGAAGCTTG GCGCCTGGG CGTGGTCTTG CAAAATGCTT CCAAAGCCAC	780
J J	CTTAGCCTGT TCTATTCAGC GGCAACCCCA AAGCACCTGT TAAGACTCCT GACCCCCAAG	840
	TOSCATOCAG CCCCCATGCC CACCOGGACC TOGTCAGCAC AGATCTTGAT GACTTCCCTT	900
40	TCTAGGGCAG ACTGGGAGGG TATCCAGGAA TCGGCCCCTG CCCCACGGGC GTTTTCATGC	960
	TGTACAGTGA CCTAAAGTTG GTAAGATGTC ATAATGGACC AGTCCATGTG ATTTCAGTAT	1020
45	ATACAACTCC ACCAGACCCC TCCAACCCAT ATAACACCCC ACCCCTGTTC GCTTCCTGTA	1080
-13	TGGTGATATC ATATGTAACA TITACTCCTG TTTCTGCTGA TTGTTTTTTT AATGTTTTGG	1140
	TTTGTTTTTG ACATCAGCTG TAATCATTCC TGTGCTGTGT TTTTTATTAC CCTTGGTAGG	1200
50	TATTAGACTT GCACTTTTTT AAAAAAAGGT TTCTGCATCG TGGAAGCATT TGACCCAGAG	1260
	TOGAACOCGT GGCCTATGCA GGTGGATTCC TTCAGGTCTT TCCTTTGGTT CTTTGAGCAT	1320
55	CTTTGCTTTC ATTOGTCTCC CGTCTTTGGT TCTCCAGTTC AAATTATTGC AAAGTAAAGG	1380
JJ	ATCTTTGAGT AGGITCGGTC TGAAAGGTGT GGCCTTTATA TTTGATCCAC ACACGTTGGT	1440
	CTTTTAACCG TGCTGAGCAG AAAACAAAAC AGGTTAAGAA GAGCCGGGTG GCAGCTGACA	1500
60	GAGGAAGCCG CTCAAATACC TTCACAATAA ATAGTGGCAA TATATATATA GTTTAAGAAG	1560

423

•	GCTCTCCATT	TGGCATCGTT	TAATTTATAT	GTTATGTTCT	AAGCACAGCT	CTCTTCTCCT	1620
5	ATTTTCATCC	TGCAAGCAAC	TCAAAATATT	TAAAATAAAG	TTTACATTGT	AGTTATTTTC	1680
J	AAATCTTTGC	TTGATAAGTA	TTAAGAAATA	TTGGACTTGC	TGCCGTAATT	TAAAGCTCTG	1740
	TTGATTTTGT	TTCCCTTTCC	ATTTTTGGGG	GAGGGGAGCA	CTGTGTTTAT	GCTGGAATAT .	1800
10	GAAGTCTGAG	ACCTTCCGGT	GCTGGGAACA	CACAAGAGTT	GTTGAAAGTT	GACAAGCAGA	1860
	CTGCGCATGT	CTCTGATGCT	TTGTATCATT	CTTGAGCAAT	CCCTCCCTCC	GTGGACAATA	1920
15	AACAGTATTA	TCAAAGAGAA	АААААААА	AAAAAACTCG	NGGGGGGCC	CGGTACCCAA	1980
13	TTCGCCCTAT	AGTGAGCCNA	TTC				2003

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(2) INFORMATION FOR SEQ ID NO: 172:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 786 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 172:

GCCACAGCGG CACGAGAAGA CTTTGGTGTT TAAGAGATTA ATGTGTTAGC CAGAACAACT 60 CATTTCTCTA COMETETETA GTCCATTTAT CTTTAAAGAT TTTCTATTGG AATAATTTTG 120 AAATTACTTT CTTAGTTTTC TTCATTAAAA ACTAAGAAAA TGCTTTGTTT ATTATGAATT 180 GCTATTTCTC TTGATTATTA TTCTTGGAGA AAGTCTATCA GACGTAATTC TTCTGATTTG 300 CTTCTAGGCT AGAGGAAAAT GTGAAAGATG ACAAATGAAA ATTTCAAAGG TTGTCAGTAG TATGACTTCT TTTATCGTTT GTCATTATCA CAAATATATC AACATAGGAC TTTTAAAAGA 420 CAGAGAGAAA GAGCAAAGAA ATAACCAAGG GTGATGTACT CGTATTGAAG GTTTACCAAA TAAGGACTGC TTTTATTATG AACTATAGTC TATATTCTAA GTAAATCAAT TTTTCTATTA 540 TGTGTTTTTT GTTCCTGCAG GCAAGATCTC TGAACTTTAT GCAGAGGGTT CTTTTAAAAA 600 AACAAAGTIG AATTITTITA TITCITGGAA TATTITTITT CATIGATITC TCCCAAGTAG 660 AGCAGATTCA AATCTCCTTT GTACCCTATG TCTTTTTTGT TTTGCTATTA GCTCAGTATT 780 ACTCGA 786

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(2) INFORMATION FOR SEQ ID NO: 173:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1758 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 173:

10	(XI) SEQUENCE DESCRIPTION. SEQ 15 NO. 1.1.	
	GOGACGAGCC CTGCCCACCT CCTGCAGCCT CCTGCGCCCC GCCGAGCTGG CGGATGGAGC	60
15	TOCGCACGGG GAGCGTGGGC AGCCAGGCGG TGGCGCGGAG GATGGATGGG GACAGCCGAG	120
15	ATGGCGGCGG CGGCAAGGAC GCCACCGGGT CGGAGGACTA CGAGAACCTG CCGACTAGCG	180
	CCTCCGTGTC CACCCACATG ACAGCAGGAG CGATGGCCGG GATCCTGGAG CACTCGGTCA	240
20	TGTACCCGGT GGACTCGGTG AAGACACGAA TGCAGAGTTT GAGTCCAGAT CCCAAAGCCC	300
	AGTACACAAG TATCTACGGA GCCCTCAAGA AAATCATGCG GACCGAAGCT TCTGGAGGCC	360
25	CTTGCGAGGC GTCAACGTCA TGATCATGGG TGCAGGGCCR GCCCATGCCA TGTATTTTGC	420
23	CTGCTATGAA AACATGAAAA GGACTTTAAA TGACGTTTTC CACCACCAAG GAAACAGCCA	480
	CCTAGCCAAC GGTATTTTGA AAGCGTTTGT CTGGAGTTAG AAAGTTCTCT TCTTCAACAC	540
30	GTCCCTCCCC AGGGTGTTCC TCCCTGTGAC CCAGCCGCCT CGACTTCGGC CCGCTTGCTC	600
	ACGAATAAAG AACTCAGAGT TGTGTGTGCA ATGCACACCC AGACACACGC ACGCACACAC	660
35	ACGCGCGCGC ACACACATGC TTTTTTCTGT TCCCCTCCGC TTTCTGAAGC CTGGGGAGAA	720
33	ATCAGTGACA GAGGTGTTTT GGTTTTATTG TTATGTGGGT TTTCTTTTGT ATTTTTTTG	780
	TTTGTTTTGT TTTTAAACAT TCAAAAGCAA TTAATGATCA GACATAGGAG AAACCCTGAA	840
40	TAGAAACAAA ACTITIGAAT GCTOGATICA AAAAAAAAA AAAGTTATCT GGACAGCTTC	900
	TTTGAGACTA TTTAAAAACT GGTACAACAG GTCTCTACAA CGCCAAGATC TAACTAAGCT	960
45	TTAAAAGGTC AAGAAGTTTT ATGGCTGACA AAGGACTCGC GCAACGCAGA AGGCCTTTCC	1020
43	CACCITAAGC TICCGGGGAT CIGGGAATTI TACCCCCATI CICITCIGIT IGICTGAGIC	1080
	TCATCTCTCT GCAAGCAAGG GCTGAAATCA TTTTGTTTGG TTGTTTTGAG GGAGAGAGGC	1140
50	GGGGTGGGGG GGTGCAAATC TGCCAGCAGC TCTTACGTAA GGCATGTTTT ATTGGGGAGG	1200
	GCTGAGCTFT TATTFTCTCC TCTCCAGTGG GGTTGGCTTT TATTGTTTCT TGTTTGGGTT	1260
55	TOGAATGGAA ATATOGATAG CAGCATAAAG TACTTTTATT TTGACAAAAT TCATTTTTT	1320
رر	CAACAATGGA GACATAGATT TGACCCACAA TAACTTCTCC CCCTCTCTTT TTACTCTGCT	1380
	CAAAAAGCAT CTCTCCTCCC ATTACCCAAC CTTGGTCATA AGTGTGCCTG GCTGGTTTGC	1440
60	ACTUAL CONTRACTOR OF THE PROPERTY OF THE PROPE	1500

	AGAGCTATGC CCTGACCTAC CCCTGATTCT ATGACATTGG GGCCCTTCTT TTGCTGAAAC	1560
5	TGCCTTACGT AATGGTTTTA CTCCTTGAAA GAGATTTGAC GGAATCCATT TTATGCCAAG	1620
•	TGCTGCCCTG CACTGTTTCT GCAATATGTG GTGTATGCTG TGGTGATCTT GCTGGGAATG	1680
	ATTATAAGTG TGTGTGGGT GGGGGAGTGG GTATTACATG CATTGCTGAA GAGTCAAAAA	1740
10	AAAAAAAAA AAACTCGA	1758
15	(2) INFORMATION FOR SEQ ID NO: 174:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 888 base pairs	
20	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 174:	
25	CTGTTAGAAT GCCCAGTTTA CCTGGATGGC AACCCAACAG TGCTCCTGCC CACCTGCCCC	60
	TCAATCCTCC TAGAATTCAG CCCCCAATTG CCCAGTTACC AATAAAAACT TGTACACCAG	120
30	CCCCAGGGAC AGTCTCAAAT GCAAATCCAC AGAGTGASMC ACCACCTCGG GTAGAATTTG	180
	ATGACAACAA TCCCTTTAGT GAAAGTTTTC AAGAACGGGA ACGTAAGGAA CGTTTACGAG	240
	AACAGCAAGA GAGACAACGG ATCCAACTCA TGCAGGAGGT AGATAGACAA AGAGCTTTGC	300
35	AGCAGAGGAT GGAAATGGAG CAGCATGGTA TGGTGGGCTC TGAGATAAGT AGTAGTAGGA	360
	CATCTGTGTC CCAGATTCCC TTCTACAGTT CCGACTTACC TTGTGATTTT ATGCAACCTC	420
40	TAGGACCCCT TCAGCAGTCT CCACAACACC AACAGCAAAT GGGGCAGGTT TTACAGCAGC	480
	AGAATATACA ACAAGGATCA ATTAATTCAC CCTCCACCCA AACTTTCATG CAGACTAATG	540
	AGCGAGGCAG GTAGGCCCTC CTTCATTTGT TCCTGATTCA CCATCAATCC CTGTTGGAAG	600
45	CCCAAATTTT TCTTCTGTGA AGCAGGGACA TGGAAATCTT TCTGGGACCA GCTTCCAGCA	660
	GICCCCAGIG AGGCCTICIT TIACACCIGC TITACCAGCA GCACCICCAG TAGCTAATAG	720
50	CAGTCTCCCA TGTGGCCAAG ATTCTACTAT AACCCATGGA CACAGTTATC CGGGATCAAC	780
	CCAATCGCTC ATTCAGTTGT ATTCTGATAT AATCCCAGAG GAAAAAGGGN AAAAAAAARA	840
	AMAARAARA ARAAAGGAGA TGATGATGCA GAATTCCACC AAGGCTCC	888
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(2) INFORMATION FOR SEQ ID NO: 175:

60 (i) SEQUENCE CHARACTERISTICS:

WO 98/54963

(A) LENGTH: 2379 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 175:

	GGCAGAGCTA	GTGTGGACTC	CATCCCCCTG	GAGTGGGATC	ACGNCTATGA	CCTCAGTCGG	. 60
10	GACCTGGAGT	CTGCAATGTC	CAGAGCTCTG	CCCTCTGAGG	ATGAAGAAGG	TCAGGATGAC	120
	AAAGATTTCT	ACCTCCGGG	AGCTGTTGSC	TTATCAGGGG	ACCACAGTGC	CCTAGAGTCA	180
15	CAGATCCGAC	AACTGGGCAA	AGCCTGGATG	ATAGCCGCTT	TCAGATACAG	CAAACCGAAA	240
13	ATATCATTCG	CAGCAAAACT	CCCACGGGGC	CGGAGCTAGA	CACCAGCTAC	AAAGGCTACA	300
	TGAAACTGCT	GGGCGAATGC	AGTAGCAGTA	TAGACTCCGT	GAAGAGACTG	GAGCACAAAC	360
20	TGAAGGAGGA	AGAGGAGAGC	CTTCCTGGCT	TTGTTAACCT	GCATAGTACC	GAAACCCAAA	420
	CCCCTCCTCT	GATTGACCGA	TGGGAGCTTC	TCCAGGCCCA	GGCATTGAGC	AAGGAGTTGA	480
25	GGATGAAGCA	GAACCTCCAG	AAGTGGCAGC	AGITTAACTC	AGACTTGAAC	AGCATCTGGG	540
23	CCTGGCTGGG	GGACACGGAG	GAGGAGTTGG	AACAGCTCCA	GCGTCTGGAA	CTCAGCACTG	600
	ACATCCAGAC	CATCGAGCTC	CAGATCAAAA	AGCTCAAGGA	GCTCCAGAAA	GCTGTGGACC	660
30	ACCGCAAAGC	CATCATCCTC	TCCATCAATC	TCTGCAGCCC	TGAGTTCACC	CAGGCTGACA	720
	GCAAGGAGAG	CCGGGACCTG	CAGGATCGCT	TGTSGCAGAT	GAATGGGCGC	TGGGACCGAG	780
35	TGTGCTCTCT	GCTGGAGGAG	TOCOCCCCC	TGCTGCAGGA	TGCCCTGATG	CAGTGCCAGG	840
<i>JJ</i>	GTTTCCATGA	AATGAGCCAT	GGTTTGCTTC	TTATGCTGGA	GAACATTGAC	AGAAGGAAAA	900
	ATGAAATTGT	CCCTATTGAT	TCTAACCTTG	ATGCAGAGAT	ACTTCAGGAC	CATCACAAAC	960
40	AGCTTATGCA	AATAAAGCAT	GAGCTGTTGG	AATCCCAACT	CAGAGTAGCC	TCTTTGCAAG	1020
	ACATGTCTTG	CCAACTACTG	GTGAATGCTG	AAGGAACAGA	CTGTTTAGAA	GCCAAAGAAA	1080
45	AAGTCCATGT	TATTGGAAAT	CGGCTCAAAC	TICTCTTGAA	GGAGGTCAGT	CGTCATATCA	1140
43	AGGAACTGGA	GAAGITATTA	GACGTGTCAA	GTAGTCAGCA	GGATTTGTCT	TCCTGGTCTT	1200
	CTGCTGATGA	ACTGGACACC	TCAGGGTCTG	TGAGTCCCAY	ATCAGGAAGG	AGCACCCCAA	1260
50	ACAGACAGAA	AACGCCACGA	GGCAAGTGTA	GTCTCTCACA	GCCTGGACCC	TCTGTCAGCA	1320
	GTCCACATAG	CAGGTCCACA	AAAGGTGGCT	CCGATTCCTC	CCTTTCTGAG	CCARGGCCAG	1380
55	GTCGGTCCGG	CCGCGGCTTC	CTGTTCAGAG	TCCTCCGAGC	AGCTCTTCCC	CTTCAGCTTC	1440
<i></i>	TCCTGCTCCT	CCTCATCGGG	CTTGCCTGCC	TTGTACCAAT	GTCAGAGGAA	GACTACAGCT	1500
	GTGCCCTCTC	CAACAACTTT	GCCCGGTCAT	TCCACCCCAT	GCTCAGATAC	ACGAATGGCC	1560
60	CTCCTCCACT	CTGAACTAAG	CAGATGCCAT	CTGCAGAAGT	GCTGGTAGCA	TAAGGAGGAT	1620

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	COGGTCATAA GCAATCCCAA ACTACCAACA AGAGGACCTT GATCTTGGCG AAAGCCMTCG	1680
5	GTGTGGCAGC TTTAGCCTCC TCCAGATCAC ATGTGTGCAA ATTATGGCTT CAGAGGTGGA	1740
J	AGATAAACAG TGACGGGGGA ACAAACAGAC AACAAGAAGG TTTGGAAGAA ATCTGGTTTG	1800
	AGACTCTGAA CCTTAGCACT AAGGAGATTG AGTAAGGACC TCCAAAGTTC CCCGGACTCA	1860
10	TGAATTCTGG GCCCTTGGCC NATTCTGTGC ACAGCCAAGG ACTTCAGTAG ACCATCTGGG	1920
	CASCITITCCC ATGGTGCTGC TCCAACCATC AGATAAATGA CCCTCCCAAG CACCATGTCA	1980
15	GTGTCGTACA ATCTACCAAC CAACCAGTGC TGAAGAGATT TTAGAACCTT GTAACATACA	2040
13	ATTITITAAGA GCTTATATGG CAGCTTCCTT TITACCTTGT TTTCCTTTGG GGCATGATGT	2100
	TITAACCTIT GCTTTAGAAG CACAAGCTGT AAATCTAAAA GGCACTTTTT TITAGAGGTA	2160
20	TAAAGAAAAA CTAGATGTAA TAAATAAGAT CATGGAAGGC TTTATGTGAA AAAAGTTGAA	2220
	TGTTATAGTA AAAAAAAAA ATATTTATGT ATGTACAGTT TGCTAAAGCC AAGTTTTGTT	2280
25	TGTATTGATT TCTTTGCATT TATTATAGAT ATTATAAAAT AAAAAAAAAA	2340
2.5	TOGAGGGGG GCCCGGTACC CAATTCGCCC TATAGTGAG	2379
30	(2) INFORMATION FOR SEQ ID NO: 176:	
	(i) SEQUENCE CHARACTERISTICS:	
35	(A) LENGTH: 1348 base pairs (B) TYPE: nucleic acid	
<i>-</i>	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 176:	
,,	GOGCCTTCAC GATGCCGGCG GTCAGTGGTC CAGGTCCCTT ATTCTGCCTT CTCCTCCTGC	60
	TCCTGGACCC CCACAGCCCT GAGACGGGGT GTCCTCCTCT ACGCAGGTTT GAGTACAAGC	120
45	TCAGCTTCAA AGGCCCAAGG CTGGCATTGC CTGGGGCTGG AATACCCTTC TGGAGCCATC	180
	ATGGAGGTGA GGGGCAGGGG TGGGGACCGC TATGCCCAGG GTCCCTCAAA GTGCTGGAGG	240
50	GGCTGTRACT TGGTGGGGAG TGGGTCTGTC ACAGCCATCC TCTGTCCAGG GTGGGGCAAG	300
J U	GCCTGGGACA GTGCCAGGCA CCCCAGGACC CCTTCCAGGC TTGTCTCCTG CTCCACCGCC	360
	TCAACACCCC CCACCCCTGC CCAAGCTGTT TCTCCTCTGC CTCTCTNNTT CCCTGCCCCA	420

55 GGACTTCTCT CTTCTCCTCT GCCTCTCCTT GGACCCCTGC CCTTCCTCTA CCTCTGACCT

60

GTGAACACAC AGACACATGC TCACACACTA AGTCCCARGC ACACMSAAAG GCAATGTGGA

CCAGCACAAA CCTCCACTCT CCCGGCTCCA TCCCARCGGG CCTGTGGCTG GCCATGAAAA

480

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	CTGGGGGCTA	CCTGGAGGGA	ACCATCCTCA	TCCCAGGTGA	GTGGGCACCA	GCCCTTCCCT	660
	GTATGTGTGT	TGTGGGTGGA	AGCAGGCATG	AGAGCATCTT	AGCCCATAGG	TTTGTATTCA	720
5	GGGACTTCCA	AACCCAGACC	TACAAAGAGT	GIGICTICTA	CCAGATCTTG	TTCAAAAAAG	780
	GGTTTGTGAT	GATGGAACTA	CACGATAGAG	GGAGTGAGCA	AGAACAATGA	GGATTAGAGT	840
10	GGAGCGTGAA	ATACTCTACG	AGCATGGCTT	CCAAAACATA	TGCTGTGAGG	TCTGTCCACC	900
10	TGAGAGTTGG	GCCATGGATT	TAATTCTGAG	CCTCTTAGCA	GCAAAGCAA	AGACAGAAAG	960
	CAGATCGGCT	GTGGATTTCT	GTCTATAAAA	TGTGAGTTCT	TOGCCCGCTG	CGGTGGCTCA	1020
15	CGCCTGTAAT	CCCGCGCTT	TGGGAGGCCA	GGCGGATGG	GTCGCGAGGT	CAGGAGGTTG	1080
	GAAACCATCC	TGGCCGGAAT	GGTGAAGCCC	TGACTCTACT	AGAAGTGCAA	AGATTGGCTG	1140
20	GGTGTGGTGG	CCTCCCCCTC	TGGTCCCAGC	TTCTCGGGAG	GCTGAGGCGG	GAGAGTTGCT	1200
20	TEECCTECC	AGGCCGAGGT	TGCGGTGAGC	TGAGATCCTG	CCATTGCACT	TCAGCCTGGG	1260
	CACAGAGCCA	GACTCTGGCT	СААААААА	АААААААА	ACTCGAGGGG	GCCCGTACC	1320
25	CAATTCGCCG	NATATGATCG	TAAACAAT				1348

30 (2) INFORMATION FOR SEQ ID NO: 177:

35

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1502 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 177:

40	CTCAAAATAA	ТАААТАААТ	AAAAATTTGT	ATTCCATTGA	TTTGGGTAGA	CACCAGGAAT	60
	GIGCATTICT	AACAAGCTTT	CCAGGCGATC	CTATAGTAAG	TCATCTGTGG	ACTACTTTAA	120
45	GAAACTCTTC	TATAGAGAAT	GGAGTTGGAT	TAATAATAGG	TGATTTTTTA	CACTGGACTG	180
73	ATTCACAAGA	ACCTAAACAG	TAGTCCATGA	AGCTGCTCAT	CTGTGGTAAC	TATTTCCCCC	240
	CGTCTCACTC	TGAAAGCAGC	aggagatgyt	GTTTACTTTG	TTTCTATCCC	CTTTGTCTGG	300
50	AGATTAATTT	TGGAATGAAA	GTTTTCTCT	CTATGCCATT	CCIGGITCIT	TTCCAAAGCC	360
	TCATACAAGA	GGATTAGGTC	ACAATGCATG	CATTACCTTT	TAAAAGAATG	CGATATTGAT	420
55	ACCGATGCTT	ACTITITIT	TTTTTNACTA	CTTGTTTTAT	TCCTTCCAGN	AAAGTATAGC	480
<i>JJ</i>	CCGCCTTTCT	ATAGCATAGT	TCTCTTTAGG	TGGAATGATT	CCTATAAGAT	TTCTCATTAT	540
	TAAATCATGC	ATTTTTCAAG	ATGGAATCAA	TMTTTGATTT	AATCTAAGCT	GATATTCTCA	600
60	TTTGTTAGAA	GAACAACCTA	CATGCTAGAG	AGAGAGGAGG	AAATATACCC	ACGACCACAC	660

429

	AGCCAGTTAG	TATCCAGTTG	GTGCTGGACT	CCAGCCAGGT	GTCCTGCCTC	ATGGTAGTTA	720
5	AATGATATAT	AGAAAAGGTA	AATTTTTAAA	GAAATATTTA	TTAATATATT	CCTATAAAAC	780
	ATTTTAAAGG	TAACCACATA	AAAATGGTTA	ATTTTTCCAT	TCCAAAGTAA	ATGCTAAGCA	840
	TGTTTATTAA	TGAAGCAGTA	CTTCTGATTA	GTATATGACA	TTCTGAAGTT	AATTAAACTC	900
10	ATTGCACTAA	ATGTGTCTTC	CTTGGTATAG	TGGAGGATTT	GAGGATTGGA	ATATAGAGTA	960
	GAGTGCTTGC	TTAAGCCTGG	GAGCCCATCT	TTATAGCTAT	TTGATGTAAG	AAAAGAGACA	1020
	TGGNCCATTT	СТАЛАСТАТА	TAAGGTGAGT	GTGTCTATTC	CCAGCAGATA	TAAAGGAAAA	1080
15	AGGAAACTTT	TTTGATTCCC	ACCTTCCCAG	CCTCACCTAG	CCATCTTCCA	GCCTCAAATA	1140
	TAGAGATGTT	AGTGCAAGGT	CCTGGGCTCT	AGGTGATCAT	TTCATAAGTC	CTTTACAGAT	1200
20	AAAGAAAAAG	TAGTGTTTGT	ATGTTTGTTT	TTAAGTAACC	CCAAAACAAA	TTTATATTGT	1260
	ATTCAGCAAA	ATTGGAATTC	AGGTGTTTAA	TTTTAGAACA	TGAAGTGCCT	GCTGTTTTAA	1320
~~	GCATTGACTT	GTATAAAAAG	AATTGCATGT	CTCCAGTAAG	CTTATGGGTT	TTCTCATTTT	1380
25	TAGGTATATG	GCTTTTAATC	ATGTAAAGTG	AAACATTAGT	TTTCTTGCAT	TTTATTACAG	1440
	GTTCTTTGTT	GCAATAAAGA	TGCTGCTGAA	AADTTAATTA	АААААААА	ааааааастс	1500
30	GA						1502

35 (2) INFORMATION FOR SEQ ID NO: 178:

40

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1637 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 178:

45 60 ATTITICTAGC CCACAAGGAC TGAAGTTCAG ATCCAAAAGT TCACTTGCTA ATTATCTTCA CAAAAATGGA GAGACTTCTC TTAAGCCAGA AGATTTTGAT TTTACTGTAC TTTCTAAAAG 120 GGGTATCAAG TCAAGATATA AAGACTGCAG CATGGCAGCC CTGACATCCC ATCTACAAAA 180 50 CCAAAGTAAC AATTCAAACT GGAACCTCAG GACCCGAAGC AAGTGCAAAA AGGATGTGTT 240 300 TATGCCGCCA AGTAGTAGTT CAGAGTTGCA GGAGAGCAGA GGACTCTCTA ACTTTACTTC 55 CACTCATTTG CTTTTGAAAG AAGATGAGGG TGTTGATGAT GTTAACTTCA GAAAGGTTAG 360 420 AAAGCCCAAA GGAAAGGTGA CTATTTTGAA AGGAATCCCA ATTAAGAAAA CTAAAAAAAGG ATGTAGGAAG AGCTGTTCAG GTTTTGTTCM AAGTGATAGC AAAAGAGAAT CTGTGTGTAA 480 60

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	TARAGCAGAT GCTGARAGTG RACCTGTTGC ACRARARAGT CAGCTTGATA GRACTGTCTG	540
	CATTTCTGAT GCTGGAGCAT GTGGTGAGAC CCTCAGTGTG ACCAGTGAAG AAAACAGCCT	600
5	TGTAAAAAAA AAAGAAAGAT CATTGAGTTC AGGATCAAAT TTTTGTTCTG AACAAAAAAC	660
	TTCTGGCATC ATAAACAAAT TTTGTTCAGC CAAAGACTCA GAACACAACG AGAAGTATGA	720
10	GGATACCTTT TTAGAATCTG AAGAAATCGG AACAAAAGTA GAAGTTGTGG AAAGGAAAGA	780
10	ACATTTGCAT ACTGACATTT TAAAACGTGG CTCTGAAATG GACAACAACT GCTCACCAAC	840
	CAGGAAAGAC TTCACTGAAG ATACCATCCC ACGGAACACA GATAGAAAGA AGGAAAACAA	900
15	GCCTGTATTT TTGCAGCAAA TATAACAAAG AAGCTCTTAG CCCCCCACGA CGTAAAGCCT	960
	TTAAGAAATG GACACCTCCT CGGTCACCTT TTAATCTCGT TCAAGAAACA CTTTTTCATG	1020
20	ATCCATGGAA GCTTCTCATC GCTACTATAT TTCTCAATCG GACCTCAGGC AAAATGGCAA	1080
20	TACCTGTGCT TTGGAAGTTT CTGGAGAAGT ATCCTTCAGC TGAGGTAGCA AGAACCGCAG	1140
	ACTOGAGAGA TOTOTCAGAA CTTCTTAAAC CTCTTGGTCT CTACGATCTT CGGGCAAAAA	1200
25	CCATTGTCAA GTTCTCAGAT GAATACCTGA CAAAGCAGTG GAAGTATCCA ATTGAGCTTC	1260
	ATGGGATTGG TGCACCCTGA AGACCACAAA TTAAATAAAT ATCATGACTG GCTTTGGGAA	1320
30	AATCATGAAA AATTAAGTCT ATCTTAAACT CTGCAGCTTT CAAGCTCATC TGTTATGCAT	1380
50	AGCTTTGCAC TTCAAAAAAG CTTAATTAAG TACAACCAAC CACCTTTCCA GCCATAGAGA	1440
	TTTTAATTAG CCCAACTAGA AGCCTAGTGT GTGTGCTTTC TTAATGTGTG TGCCAATGGT	1500
35	GGATCTTTGC TACTGAATGT GTTTGAACAT GTTTTGAGAT TTTTTTAAAA TAAATTATTA	1560
	ТТТСАСЛАСА АТССАЛЛАЛА АЛЛДАЛАЛА ЛАЛДАЛАЛА ЛАЛДАЛАЛА АЛЛДАЛАЛА	1620
40	AAAAAAA AAAAAAA	1637
70		
45	(2) INFORMATION FOR SEQ ID NO: 179:	
45	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 2911 base pairs (B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	•
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 179:	
<i></i>	GGTGGTTTTT GTTCTGCAAT AGGCGGCTTA GAGGGAGGGG CTTTTTCGCC TATACCTACT	60
55	GTAGCTTCTC CACGTATGGA CCCTAAAGGC TACTGCTGCT ACTACGGGGC TAGACAGTTA	120
	CTGTCTCAGC TCTAGGATGT GCGTTCTTCC ACTAGAAGCT CTTCTGAGGG AGGTAATTAA	180
60	AAAACAGTGG AATGGAAAAA CAGTGCTGTA GTCATCCTGT AATATGCTCC TTGTCAACAA	240

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	TGTATACATT CCTGCTAGGT GCCATATTCA TTGCTTTAAG CTCAAGTCGC ATCTTACTAG	300
c	TGAAGTATTC TGCCAATGAA GAAAACAAGT ATGATTATCT TCCAACTACT GTGAATGTGT	360
5	GCTCAGAACT GGTGAAGCTA GTTTTCTGTG TGCTTGTGTC ATTCTGTGTT ATAAAGAAAG	420
	ATCATCAAAG TAGAAATTTG AAATATGCTT CCTGGAAGGA ATTCTCTGAT TTCATGAAGT	480
10	GGTCCATTCC TGCCTTTCTT TATTTCCTGG ATAACTTGAT TGTCTTCTAT GTCCTGTCCT	540
	ATCTTCAACC AGCCATGGCT GTTATCTTCT CAAATTTTAG CATTATAACA ACAGCTCTTC	600
٠, ٣	TATTCAGGAT AGTGCTGAAG ANGCGTCTAA ACTGGATCCA GTGGGCTTCC CTCCTGACTT	660
15	TATTTTTGTC TATTGTGGCC TIGACTGCCG GGACTAAAAC TTTACAGCAC AACTTGGCAG	720
	GACGTGGATT TCATCACGAT GCCTTTTTCA GCCCTTCCAA TICCTGCCTT CTTTTCAGAA	780
20	ATGAGTGTCC CAGAAAAGAC AATTGTACAG CAAAGGAATG GACTTTTCCT GAAGCTAAAT	840
	GGAACACCAC AGCCAGAGTT TTCAGTCACA TCCGTCTTGG CATGGGCCAT GTTCTTATTA	900
25	TAGTCCAGTG TTTTATTTCT TCAATGGCTA ATATCTATAA TGAAAAGATA CTGAAGGAAG	960
25	GGAACCAGCT CACTGAARGC ATCTTCATAC AGAACAGCAA ACTCTATTTC TTTGGCATTC	1020
	TGTTTAATGG GCTGACTCTG GGCCTTCAGA GGAGTAACCG TGATCAGATT AAGAACTGTG	1080
30	GATTTTTTA TOGCCACAGT GCATTTTCAG TAGCCCTTAT TTTTGTAACT GCATTCCAGG	1140
	GCCTTTCAGT GGCTTTCATT CTGAAGTTCC TGGATAACAT GTTCCATGTC TTGATGGCCC	1200
25	AGGITACCAC TGTCATTATC ACAACAGTGT CTGTCCTGGT CTTTGACTTC AGGCCCTCCC	1260
35	TOGAATTTTT CTTGGAAGCC CCATCAGTCC TTCTCTCTAT ATTTATTTAT AATGCCAGCA	1320
	AGCCTCAAGT TCCGGAATAC GCACCTAGGC AAGAAAGGAT CCGAGATCTA AGTGGCAATC	1380
40	TTTGGGAGCG TTCCAGTGGG GATGGAGAAG AACTAGAAAG ACTTACCAAA CCCAAGAGTG	1440
	ATGAGTCAGA TGAAGATACT TTCTAACTGG TACCCACATA GTTTGCAGCT CTCTTGAACC	1500
45	TTATTITCAC ATTITCAGIG TITGIAATAT TTATCTITTC ACTITGATAA ACCAGAAATG	1560
45	TTTCTAAATC CTAATATTCT TTGCATATAT CTAGCTACTC CCTAAATGGT TCCATCCAAG	1620
	GCTTAGAGTA CCCAAAGGCT AAGAAATTCT AAAGAACTGA TACAGGAGTA ACAATATGAA	1680
50	GAATTCATTA ATATCTCAGT ACTTGATAAA TCAGAAAGTT ATATGTGCAG ATTATTTTCC	1740
	TTGGCCTTCA AGCTTCCAAA AAACTTGTAA TAATCATGTT AGCTATAGCT TGTATATACA	1800
	CATAGAGATC AATTTGCCAA ATATTCACAA TCATGTAGTT CTAGTTTACA TGCCAAAGTC	1860
55	TICCCTTTT AACATTATAA AAGCTAGGIT GICTCTIGAA TTTIGAGGCC CTAGAGATAG	192
	TCATTTTGCA AGTANAGAGC AACGGGACCC TITCTAANAA CGTTGGTTGA AGGACCTANA	198
60	TACCTGGCCA TACCATAGAT TTGGGATGAT GTAGTCTGTG CTAAATATIT TGCTGAAGAA	204

	GCAGTTTCTC AGACACAACA TCTCAGAATT TTAATTTTTA GAAATTCATG GGAAATTGGA	2100
_	TTTTTGTAAT AATCTTTTGA TGTTTTAAAC ATTGGTTCCC TAGTCACCAT AGTTACCACT	2160
5	TGTATTTTAA GTCATTTAAA CAAGCCACGG TGGGGCTTTT TTCTCCTCAG TTTGAGGAGA	2220
	AAAATCTTGA TGTCATTACT CCTGAATTAT TACATTTTGG AGAATAAGAG GGCATTTTAT	2280
10	TITATTAGIT ACTAATICAA GCTGTGACTA TIGTATATCT TICCAAGAGI TGAAATGCTG	2340
	GCTTCAGAAT CATACCAGAT TGTCAGTGAA GCTGATGCCT AGGAACTTTT AAAGGGATCC	2400
	TTTCAAAAGG ATCACTTAGC AAACACATGT TGACTTTTAA CTGATGTATG AATATTAATA	2460
15	CTCTAAAAAT AGAAAGACCA GTAATATATA AGTCACTTTA CAGTGCTACT TCACACTTAA	2520
	AAGTGCATGG TATTTTCAT GGTATTTTGC ATGCAGCCAG TTAACTCTCG TAGATAGAGA	2580
20	AGTCAGGTGA TAGATGATAT TAAAAATTAG CAAACAAAAG TGACTTGCTC AGGGTCATGC	2640
	ACCTGGGTGA TGATAGAAGA GTGGGCTTTA ACTGGCAGGC CTGTATGTTT ACAGACTACC	2700
05	ATACTGTAAA TATGAGCTTT ATGGTGTCAT TCTCAGAAAC TTATACATTT CTGCTCTCCT	2760
25	TTCTCCTAAG TTTCATGCAG ATGAATATAA GGTAATATAC TATTATATAA TTCATTTGTG	2820
•	ATATCCACAA TAATATGACT GGCAAGAATT GGTGGAAATT TGTAATTAAA ATAATTATTA	2880
30	аасставава ававававав ававастсда д	2911

35 (2) INFORMATION FOR SEQ ID NO: 180:

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60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 519 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 180:

45	GGCACGAGCC CCAGGCCAGC CAGGGCCAGG CCTACTTTGG CCACCCTTAA ATTAGAATGT	60
	GGGTCAGGG GTCACAGAAA AGCCATTTCT CTGACCTAGT GTTTGGCGTC CGGGAACTCT	120
50	GTGCCCAACC TTCAGACCCT GGCAGTCCTC ACTGAGGCCA TTGGCCCAGA GCCCGCCATC	180
50	CCCCGARACC CCCGGGAGCC GCCTGTTGCC ACGTCCACAC CTGCCCACACC CTCTGCCGGG	240
	CCCCAGCCCC TCCCAACCGG GACCGTGCTG GTCCCTGGGG GTCCTGCCCC ACCTTGCCTT	300
55	GGGGAGGCAT GGGCCCTCCT CCTCCCACCC TGCCGGCCGT CACTCACCTC TTGCTTCTGG	360
	TOCCCCAGGC CTAGCCCTTG GAAGGAGACA GGAGTCTAGG GAGGCTGAAG CCCACTCCCG	420
	GGGAGGCCCG TGCTCCTCCA GCCCCAGGGA CAGCAAGGAA AAGAGAAGAG	480

433

TTCATGGCTC TAATAAAAA	AAAAAAAAA AAAACTCGA	519
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5
(2) INFORMATION FOR SEQ ID NO: 181:

10

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 968 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 181:

15 TCCCCTTGGG GCCGGAAAAA GCGGGGTTGG CCTGNCCATT GGTTNTCCAT GCCGCCCGCC 60 CATGCCCCAG TACTAGCCTG CAGTCCCAAT GTAGCCCCTC CCTCYTCCMA GAGCCCYTCM 120 AACCGCCCCG STCANTIGIG ATTICAGGAG GATTIGATGA AGATGITAAA GCGAAAGTGG 180 20 AGAACCTTCT CGGGATTTCC AGCCTGGAAA AAACGGACCC TGTTAGGCAA GCACCCTGCA 240 GCCCTCCCTG TCCCCTTCTT CCCCTCCCCT TCYCCCGCCC GTGGAGACAG CTGTTYTCAG 300 25 360 CAGGGCTCTC CGCAGGGAGG GGGCCGGCTC CTTCCCTGGC AGCAACATCC TTGCCCTTGT CACACAAGTC AGCCTCCATC TGCGCAGCTC TGTGGATGCG CTGCTGGAGG GCAACAGGTA 420 TGTCACTGGC TGGTTCAGCC CCTACCACCG CCAGCGGAAG CTCATCCACC CGGTCATGGT 480 30 TCAGCACATC CAGCCCGCAG CGCTCAGCCT CCTGGCACAG TGGAGCACCC TCGTGCAGGA 540 GCTGGAGGCT GCCCTGCAGC TGGCTTTCTA CCCGGATGCC GTGGAGGAGT GGCTGGAGGA 600 35 AAACGTGCAC CCCAGCCTGC AGCGGCTGCA ARCTCTGCTG CAGGACCTCA GCGAGGTGTC 660 TGCCCCCCG CTGCCACCCA CCAGCCCTGG CAGGGACGTT GCTCAGGACC CCTGAGGGGA 720 780 GAGCTCATGC CAGGGGGCTC CTGCTGGAGG CTGGGGGGGC TCTGCWYTKY CWWWTGGCCT 40 GGGCAATACG GCCCACGTGG GCGTCGTGCC CTCTGGCCCA GCAGTGTCTT GCCCACACTC 840 AGTTCCTGAG GGCCCTGGGC AGCCCCTGGG GGAGAGACTA GAAAACACAG AAGGAAGCAG 900 45 CACAGGGAGA CCCGCTTTGT GATCTGCATG TGTGACACTG ATTCTTTGGA AATAAAGAGT 960 968 GGAAGCTG

(2) INFORMATION FOR SEQ ID NO: 182:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1128 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

60

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 182:	
	TGTAAAAGTT ATCAGTAATC CTAATTCTTT TCCTGGGTTT TCCTTTTGTC ACTTATTAAT	60
5	CAGTITITIGA AAGGACGAAT GAATITAGAG ATGTACTCTG GAGCAGTATC ATGTTAAACC	120
	AGGGGTATAT TAGAAAAATC ATCCTCATAA TCATTCTGGG AAGTTTTTCC TCCCCAAAAA	180
	AAGCCATCCT GATGGGTTTT CAAAACCAGA AAAAAGCTCT TAATGAGGAA CAGACCACTG	240
10	GAGTACCCAT GAGCATCTCA GGAAAACTGA GACCCTCGAG AAGCCTTGAT TTCGTGCAAC	300
	CCCCAAGGIT TCAGAGCCAG CAGCCCAGTG CTGTGGTTGA CAGACGTGGT TTTKTGGRGA	360
15	AAGCAGCCAG AGGCCAGGAA TTTTCAGAGT CGTGAGTCAC GRTYTCCCAC CCAAGATTAG	420
	AGCAMAGATT AGCCATACTG AGATTTGGTA AAATCATTCT GTCTAAGCAA TGGAGGTGTG	480
	TOCAMACOTO CAGTOCCTOT TCACAGOGGA TGCAGGCAGA TCSYGGOTTT AGGATGGGGR	540
20	AGGCCACCGC ACCCCCYTTC AYTGCTCTGC ACCTGCTCCC TCACGTGGAC ACTGTCCACA	600
	ACTIGTOSCTC TCACAGGACA GTTGCCCCAAG GAGCTCATAT CTTATTGGAG ATAGGGGGTC	660
25	GTACAGGTGA CATTCATGAG CAGTGTGAGC CGGGTGACAT GGGGGTGTCA ACCCAGCATC	720
	TGTCCAGGAG CTCCTCCTGC AGCGGCTCTG GCAGGTGGCC TGAGGCTCCT TTTTGAGAGA	780
20	GAACTGTTTG GCCTTCCTGT CTCCTCTCCT CTGATCTGTT CTTTCTTGGA ACACCACCCA	840
30	AGAACGTCAC CTCCTCCATC AGATTGTGAG CTCCTGGAGG GCAGGAGCTG TGTCCTTCTA	900
	TTCATCTTCC TATCCCCAGA ACCTTGCACA GATCCTGGAA TGTGGTAGGT GCTCAGTAAA	960
35	TGTGTGTGA ATAAATGAAT GAATGAATGA ACAAATGAAT GAATTTGCTT ACTTCAAGGC	1020
	AAAAGAACCA TGAAACTGTA TTTTGAGTTT CTATGTTATA GCAGTCAGCA AATCCTATTA	1080
40	AATACTTIGT GITTCCAAGC AAAAAAAAAA AAAAAAAAA AAACTCGA	1128
40		
45	(2) INFORMATION FOR SEQ ID NO: 183:	
.5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2276 base pairs	
	(B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 183:	
	COGCOGCOTO TGACOTOATO GOOTAGAGOO TAGCAACAGO GCAGGOTOCO AGCOGAGTOO	60
55	GTTATGGCCG CTGCCGTCCC GAAGAGGATG AGGGGGCCAG CACAAGCGAA ACTGCTGCCC	120
		180
	GGGTCGGCCA TCCAAGCCCT TGTGGGGTTG GCGCGGCCGC TGGTCTTGGC GCTCCTGCTT	
60	GTGTCCGCCG CTCTATCCAG TGTTGTATCA CGGACTGATT CACCGAGCCC AACCGTACTC	240

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	AACTCACATA TTTCTACCCC AAATGTGAAT GCTTTAACAC ATGAAAAACCA AACCAAACCT	300
_	TCTATTTCCC AAATCAGCAC CACCCTCCCT CCCACGACGA GTACCAAGAA AAGTGGAGGA	360
5	GCATCTGTGG TCCCTCATCC CTCGCCTACT CCTCTGTCTC AAGAGGAAGC TGATAACAAT	420
	GAAGATCCTA GTATAGAGGA GGAGGATCTT CTCATGCTGA ACAGTTCTCC ATCCACAGCC	480
10	AAAGACACTC TAGACAATGG CGATTATGGA GAACCAGACT ATGACTGGAC CACGGCCCC	540
	AGGGACGACG ACGAGTCTGA TGACACCTTG GAAGAAAACA GGGGTTACAT GGAAATTGAA	600
	CAGTCAGTGA AATCTTTTAA GATGCCATCC TCAAATATAG AAGAGGAAGA CAGCCATTTC	660
15	TTTTTTCATC TTATTATTTT TGCTTTTTGC ATTGCTGTTG TTTACATTAC ATATCACAAC	720
	AAAAGGAAGA TTTTTCTTCT GGTTCAAAGC AGGAAATGGC GTGATGGCCT TTGTTCCAAA	780
20	ACAGTGGAAT ACCATCGCCT AGATCAGAAT GITAATGAGG CAATGCCTTC TTTGAAGATT	840
	ACCAATGATT ATATTTTTA AAGCACTGTG ATTTGAATTT GCTTATGTAA TTTTATTTGC	900
	TIGACTITIT ATATGATATT GIGCAAATGI TIGCCATAGG CAATIGGIAC TIAAATGAGA	960
25	GGTGAGTCTC TCTTTTGCCT TGGTGCTTTG GAAATTAAAT GTCACAAACG AGTATATAAT	1020
	TTTTTATCTG TACTTTTAGA GCTGAGTTTA ATCAGGTGTC CAAAATGTGA GTTAAACATT	1080
30	ACCITATATI TACACIGITA GITTITATIG TITTAGATIT ATTAIGCTIC TICIGGAAGI	1140
	ATTAGTGATG CTACTTTTAA AAGATCCCAA ACTTGTAACT AAATTCTGAC ATATCTGTTA	1200
	CTGCTGACTC ACATTCATTC TCCGCCATTC AAATACTATT TITTATCCAC ATTTTTTTTT	1260
35	GTTCCCAAAC TGTAATGTAC AAGGATATGT GTGATAATGC TTTGGATTTG AGTAATATTT	1320
	TTTTTCTTC CAAGAAAACT GCTTTGGATA TTTTTAGATA ATTTAAACAT AATTTAGGAT	1380
40	AATGATATTG CTCAATCTGA CCACAATTTT AGGTAAAACA TTAAATGTGT CAGAAATCTT	1440
	GGCAACAGAG ACTCTGCAGC TTGCAGTGGA CATAGATAAA ATGTTACAGA GATACTATTT	1500
45	TTTTGGTTGG AATTACTATA TTAAATTTAG AAGCAGAAAC TGGTAAAATG TTAAATACAT	1560
45	GTACAATIGC TITTAGTTAG CAATIGATIG TAGCATGGGT TCCTCCAAGG TITCAAGCAA	1620
	TOGGCAGAGT TTAAAATTAT ATCAGATTCG TITACTTCGT TTATTATTTT ACAGTAAATT	1680
50	TGAATAAATC TTAGGGGTCA TTATCACTTA AATAATACTG TACCTAGGTC TTTCAAATTA	1740
	AAATTATACC TGAATGAAGT TGTTTGTATA CATAAAGGAT ATTTGTGTAC AATTACCTTT	1800
	TTTCCCCCAC ACTIGITITIC TITIGITTITG TTTTTTATGG CAACTGGAAA GTATTTACTA	1860
55	TGGGATTCAT TTATGTCTGT CTTTCTATCA TAAAGAATTG ATCAATATGT AAATATGTGA	1920
	TTTGAACCAT GGTTGACTTA CAAGTGTCAC TACAGCTTTT TAGAAAACAT AGCCCTAATA	1980
60) TATGTTAACC AGGACCCGGG TGAGCCAGTG GGCTTGCGCT TTATGTAGAG CTGGAAGAAG	2040

	GCCGTCCATC CTGTCTCTTG GGCGGACAGT GTACTTTCCT AATAGGGAAG GGAAGCACAA	2100
5	TOGANATACC CCTGAACCGT TTTATTGCAG TAATTTTTTT CATATCTGAA ACTATTATTT	2160
	AATATTTTGA ATAAGATTT AAAAAATAAA TOOCAAAGAT ATAAATCTAA AAAAAAAAA	2220
	AAAAAA AAAAAAAAA AAAAAAAAA AAAAAAAAA AAAA	2276
10		
10		
	(2) INFORMATION FOR SEQ ID NO: 184:	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2500 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
20		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 184:	60
	TCCAAGCTAC GCCACTCGGG CTGGGGGCGTT GGGAGCGGGA GTGCAGAGCG TGGTCGTGGC	60
25	GGCGGCGGTG AGAAGAGCGA GGCGKAGGAG GGGGTGCCAT GGCCGGGCAG CAGTTCCAGT	120
	ACGATGACAG TGGGAACACC TTCTTCTACT TCCTCACCTC CTTCGTGGGG CTCATCGTGA	180
	TCCCGGCGAC ATACTACCTC TGGCCCCGAG ATCAGAATGC CGAGCAAATT CGATTAAAGA	240
30	ATATCAGAAA AGTATATGGA AGGTGTATGT GGTACGTTTA CGGTTATTAA AACCCCAGCC	300
	AAATATTATT CCTACAGTAA AGAAAATAGT TCTGCTTGCA GGATGGGCAT TGTTCTTATT	360
35	CCTTGCATAT AAAGTTTCCA AAACAGACCG AGAATACCAA GAATACAATC CTTATGAAGT	420
	ATTARATTIG GATCCIGGAG CCACAGTAGC AGRARTTARA ARACARTRIC GITTGCIGIC	480
	ACTTAAATAT CATCCAGATA AAGGAGGTGA TGAGGTTATG TTCATGAGGA TAGCAAAAGC	540
40	TTATGCTGCT TTAACGGATG AAGAGTCCCG GAAAAATTGG GAAGAATTTG GAAATCCAGA	600
	TGGCCTCAA GCCACAAGCT TTGGAATTGC CCTGCCAGCT TGGATAGTTG ACCAGAAAAA	660
45		720
	GGGCTCTTGG TGGTATCGCT CAATACGCTA TAGTGGAGAC CAGATTCTAA TACGSACAAC	780
	ACAGATITAT ACATACTITG TITATAAAAC CCGAAATATG GATATGAAAC GTCTTATCAT	840
50		900
	AACGGATAAT ATTCTAATAC CACAGCTAAT CAGAGAAATT GGCAGCATTA ATTTAAAGAA	960
<<		1020
55	TCTTGCTAGA ATGAAAATTC CTGAGACCCT TGAAGAAGAT CAGCAATTCA TGCTAAAAAA	
	•	114
	GTGTCCTGCC CTACTTCAAG AAATGGTTAA TGTAATCTGC CAACTAATAG TAATGGCCCG	

	GAACCGTGAA GAAAGGGAGT TTCGTGCTCC AACTTTGGCA TCCCTAGAAA ACTGCATGAA	1200
	GCTTTCTCAG ATGGCCGTTC AGGGACTTCA GCAATTTAAG TCTCCCCTTC TGCAGCTCCC	1260
5	TCATATTGAA GAGGACAATC TTAGACGGGT TTCTAATCAT AAGAAGTATA AAATTAAAAC	1320
	TATCCAGGAT TIGGTGAGTT TAAAAGAATC AGATCGTCAC ACTCTACTGC ACTTCCTTGA	1380
10	AGATGAAAAA TATGAAGAGG TTATGGCTGT CCTTGGGAGT TTTCCATATG TGACCATGGA	. 1440
10	TATAAAATCA CAGGTGTTAG ATGATGAAGA TAGCAACAAC ATCACAGTAG GATCCTTAGT	1500
	TACAGTGTTG GTTAAGTTGA CAAGGCAAAC AATGGCTGAA GTATTTGAAA AGGAGCAGTC	1560
15	CATCTGTGCT GCAGAGGAAC AGCCAGCAGA AGATGGGCAG GGTGAAACTA ACAAGAACAG	1620
	GACAAAAGGA GGATGGCAAC AGAAGAGTAA AGGACCCAAG AAAACTGCTA AATCAAAAAA	1680
20	AAAGAAACCT TTAAAAAAAA AACCTACACC TGTGCTATTA CCACAGTCAA AGCAACAGAA	1740
20	ACAAAAGCAG GCAAATGGAG TCGTTGGGAA TGAAGCTGCA GTAAAGGAAG ATGAAGAAGA	1800
	AGTITICAGAT AAGGGCAGTG ATTCTGAAGA AGAAGAAACC AATAGAGATT CCCAAAGTGA	1860
25	GAAAGATGAT GGTAGTGACA GAGACTCTGA TAGAGAGCAA GATGAAAAAC AAAACAAAGA	1920
	TGATGAAGCA GAGTGGCAAG AATTACAACA AAGCATACAG CGAAAAGAGA GAGCTCTATT	1980
30	GGAAACCAAA TCAAAAATAA CACATCCTGT GTATAGCCTT TACTTTCCTG AGGAAAAACA	2040
30	AGAATGGTGG TGGCTTTACA TTGCAGATAG GAAGGAGCAG ACATTAATAT CCATGCCATA	2100
	TCATGTGTGT ACGCTGAAAG ATACAGAGGA GGTAGAGCTG AAGTTTCCTG CACCAGGCAA	2160
35	GCCTGGAAAT TATCAGTATA CTGTGTTTCT GAGATCAGAC TCCTATATGG GTTTGGATCA	2220
	GATTAAACCA TTGGAAGTTK GGAAGTTCAT GAGGCTGAAG CCTGTGCCAG AAAATCACCC	228
40	ACAGTGGGAT ACAGCAATAG AGGGGGATGA AGACCAGGAG GACAGTGAGG GCTTTGAAGA	234
40	TACCTTTGAG GGAGGAAGAG GGAGGGAGGA AGGAAGGTGG TGGACTTAAG GCAGTTACTC	240
	TOGANTOGON COCACAGIGT TITTGCACCAT ATTITTGGCAN TITTITTTTGC CCGITTTING	246
45	GAACTICTTTT CONTINANCO CAGGAACCAT TACAGAACCG	250

50 (2) INFORMATION FOR SEQ ID NO: 185:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1337 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 185:

60 CTTCCGGTTC TCCGGGCAGC TGCCACTGCT GTAGCTTCTG CCACCTGCCA CGACCGGGCC

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	TCTCCCTGGC GTTTGGTCAC CTCTGCTTCA TTCTCCACCG CGCCTATGGT CCCTCTTGGA	120
	GCCAGCGTGG CGGGCCTGGC GGCTCCCGGG TGGTGAGAGA GCGGTCCGGG AACGATGAAG	180
5	GCCTCGCAGT GCTGCTGCTG TCTCAGCCAC CTCTTGGCTT CCGTCCTCCT CCTGCTGTTG	240
	CTGCCTGAAC TAAGCGGGYC CCTGGMAGTC CTGCTGCAGG CAGCCGAGGC CGCGCCAGGT	300
10	CTTGGGCCTC CTGACCCTAG ACCACGGACA TTACCGCCGC TGCCACCGGG CCCTACCCCT	360
	GCCCAGCAGC CGGGCCGTGG TCTGGCTGAA GCTGCGGGGC CGCGGGGCTC CGAGGGAGGC	420
	AATGGCAGCA ACCCTGTGGC CGGGCTTGAG ACGGACGATC ACGGAGGGAA GGCCGGGGAA	480
15	GOCTOGGTGG GTGGCGGCCT TGCTGTGAGC CCCAACCCTG GCGACAAGCC CATGACCCAG	540
	CGGGCCCTGA CCGTGTTGAT GGTGGTGAGC GGCGCGGTGC TGGTGTACTT CGTGGTCAGG	600
20	ACGGTCAGGA TGAGAAGAAG AAACCGAAAG ACTAGGAGAT ATGGAGTTTT GGACACTAAC	660
	ATAGAAAATA TGGAATTGAC ACCTTTAGAA CAGGATGATG AGGATGATGA CAACACGTTG	720
	TTTGATGCCA ATCATCCTCG AAGATAAGAA TGTGCCTTTT GATGAAAGAA CTTTATCTTT	780
25	CTACAATGAA GAGTGGAATT TCTATGTTTA AGGAATAAGA AGCCACTATA TCAATGTTGG	840
	GGGGTATTT AAGTTACATA TATTTTAACA ACCTTTAATT TGCTGTTGCA ATAAATACCG	900
30	TATCCTTTTA TTATATCTTT ATATGTATAG AAGTACTCTR TTAATGGGCT CAGAGATGTT	960
	GGGGATAAAG TATACTGTAA TAATTTATCT GTTTGAAAAT TACTATAAAA CGGTGTTTTC	1020
25	TGATCGGTTT TTGTTTCCTG CTTACCATAT GATTGTAAAT TGTTTTATGT ATTAATCAGT	1080
35	TAATGCTAAT TATTTTTGCT GATGTCATAT GTTAAAGAGC TATAAATTCC AACAACCAAC	1140
	TGGTGTGTAA AAATAATTTA AAATTTCCTT TACTGAAAGG TATTTCCCAT TTTTGTGGGG	1200
40	AAAAGAAGCC AAATITATTA CTTTGTGTTG GGGTTTTTAA AATATTAAGA AATGTCTAAG	1260
•	TTATTGTTTG CAAAACAATA AATATGATTT TAAATTCTCT TAAAAAAAAA AAAAAAAACC	1320
15	CCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCC	1337

(2) INFORMATION FOR SEQ ID NO: 186:

50
(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 941 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 186:

GGCACGAGCC TGGACGCAGC AGCCACCGCC GCGTCCCTCT CTCCACGAGG CTGCCGGCTT 60

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	AGGACCCCCA GCTCCGACAT GTCGCCCTCT GGTCGCCTGT GTCTTCTCAC CATCGTTGGC	120
	CTGATTCTCC CCACCAGAGG ACAGACGTTG AAAGATACCA CGTCCAGTTC TTCAGCAGAC	180
5	TCAACTATCA TOGACATTCA GGTCCCGACA CGAGCCCCAG ATGCAGTCTA CACAGAACTC	240
	CAGCCCACCT CTCCAACCCC AACCTGGCCT GCTGATGAAA CACCACAACC CCAGACCCAG '	300
10	ACCCAGCAAC TGGAAGGAAC GGATGGGCCT CTAGTGACAG ATCCAGAGAC ACACAAGAGC .	360
10	ACCAAAGCAG CTCATCCCAC TGATGACACC ACGACGCTCT CTGAGAGACC ATCCCCAAGC	420
	ACAGACGTCC AGACAGACCC CCAGACCCTC AAGCCATCTG GTTTTCATGA GGATGACCCC	480
15	TTCTTCTATG ATGAACACAC CCTCCGGAAA CGGGGGCTGT TGGTCGCAGC TGTGCTGTTC	540
	ATCACAGGCA TCATCATCCT CACCAGTGGC AAGTGCAGGC AGCTGTCCCG GTTATGCCGG	600
20	AATCATTIGCA GGTGAGTCCA TCAGAAACAG GAGCTGACAA CCYGCTGGGC ACCCGAAGAC	660
20	CAAGCCCCCT GCCAGCTCAC CGTGCCCAGC CTCCTGCATC CCCTCGAAGA GCCTGGCCAG	720
	AGAGGGAAGA CACAGATGAT GAAGCTGGAG CCAGGGCTGC CGGTCCGAGT CTCCTACCTC	780
25	CCCCAACCCT GCCCGCCCCT GAAGGCTACC TGGCGCCTTG GGGGCTGTCC CTCAAGTTAT	840
	CTCCTCTGYT AAGACAAAAA GTAAAGCACT GTGGTCTTTG CAAAAAAAAAA	900
30	алалалала алалалала алалалала алалаластСG A	941
35	(2) INFORMATION FOR SEQ ID NO: 187:	
33	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 654 base pairs	
	(B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 187:

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GAATTCGGCA CGAGGCAGCT TGTGCTTTAA AGGAGGTGTT CAAAGCATGT CTGAGCAGAG 60 ACTITIGGG TCIGITITAA TTAATACTTI AAAATAATTC ATATITAAAA TATCARATGI TTCCATAAAG AGGAGGATGT TTAAATGCCT CCAGACTACA TTCCTTTTTA TTSCTTGATT 180 TTACCTOGGA GTCCAAAGTT CAATTCCCAT AAAGCAAGCG TTTTATTTGT CACTTTCAAT 240 ATACATCCGA TTGCCATGCT TAAGATGCAA TATGGGCTGC GGAAATAGGT TAACCCACAG 300 GCTCCCAGGG CCCAGTGTAG AAGGTGAGAG ATTCGTGTAA AATGATTCAA ATAAAAGGAA 360 GACCCTGGCC GGGTGCCGTA RCTCACGCCT GTAATCCCAG CACTTTGGGA GGCCGAAGCG 420 ACTGGATGAC GAGGTTAGGA GTTGGAGACC AGCCTGGCCA ACATCGTGAA ACCCCGTCTC 480 TACTAAAAAT ACAAAAATTA GCCGGGCATG GTGGCAGGCA CCTGTAATCC TAGCTAGTTG 540

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	GGAGGCTGAG GCAGGAGAAT CGTTTGAATC TGGGAGTTGG AGGTTGTCAG TGAGCTGAGA	600
5	TCGCGCCACA GCACTCCAGC CTGGGTGACA GGGTGAGACT CTGTCTCAAA NAGA	654
10	(2) INFORMATION FOR SEQ ID NO: 188: (i) SEQUENCE CHARACTERISTICS:	
15	(A) LENGTH: 1848 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 188:	
20	GAAACTOGAC COGAGAACCG GAGCGAAGCG AAGCGGAAGC CCGGAATGAG GCCGGACTGG	60
20	AAAGCCGGAG CGGGCCAGG CGGGCCTCCC CAAAAGCCTG CCCCTTCATC CCAGCGGAAA	120
	CCGCCGGCCC GGCCGAGCGC GGCGGCCGCT GCGATTGCAG TCGCGGCGGC GGAGGAAGAG	180
25	AGACGGCTCC GGCAGCGGAA CCGCCTGAGG CTGGAGGAGG ACAAACCGGC CGTGGAGCGG	240
	TECTTEGAGG ACCTEGTCTT CECCGACCTC GAGAACGACG AGGACGCCTT GCTCCGGCCT	300
20	CTGCGAGGCC CGAGGGTTCA AGAACATGAA GACTCGGGTG ACTCAGAAGT GGAGAATGAA	360
30	GCAAAAGGTA ATTTTCCACC TCAAAAGAAG CCAGTTTGGG TGGATGAAGA AGATGAAGAT	420
	GAGGAAATGG TTGACATGAT GAACAATCGG TTTCGGAAGG ATATGATGAA AAATGCTAGT	480
35	GAAAGTAAAC TITCGAAAGA CAACCTTAAA AAGAGACTTA AAGAAGAATT CCAACATGCC	540
	ATGGGAGGAG TACCTGCCTG GGCAGAGACT ACTAAGCGGA AAACATCTTC AGATGATGAA	600
40	AGTGAAGAGG ATGAAGATGA TITGTTGCAA AGGACTGGGA ATTTCATATC CACATCAACT	660
40	TCTCTTCCAA GAGGCATCTT GAAGATGAAG AACTGCCAGC ATGCGAATGC TGAACGTCCT	72 0
•	ACTOTTOCTC GGATCTCCAT CTGTGCAGTT CCATCCCGGT GCACAGATTG TGATGGTTGC	780
45	TGGGATTAGA TAATGCTGTA TCACTATTTC AGGTTGATGG GAAAACAAAT CCTAAAATTC	840
	AGAGCATCTA TTTGGAAAGG TTTCCAATCT TTAAGGCTTG TTTTAGTGCT AATGGGGAAG	900
	AAGTTTTAGC CACGAGTACC CACAGCAAGG TTCTTTATGT CTATGACATG CTGGCTGGAA	960
50	AGTTAATTCC TGTGCATCAA GTGAGAGGTT TGAAAGAGAA GATAGTGAGG AGCTTTGAAG	1020
	TCTCCCCAGA TGGGTCCTTC TTGCTCATAA ATGGCATTGC TGGATATTTG CATTTGCTAG	1080
55	CAATGAAGAC CAAAGAACTG ATTGGAAGCA TGAAAATTAA TGGAAGGGTT GCAGCATCCA	1140
	CATTCTCTC AGATAGTAAG AAAGTATACG CCTCTTCGGG GGATGGAGAA GTTTATGTTT	1200
	THE TAXABLE PROPERTY OF COMMENCES WELLOCOLOGY TO THE TRANSPORT	1260

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TAAGCATTGC CACATCTAGG AATGGACAGT ATGTTGCTTG TGGTTCTAAT TGTGGAGTGG	1320
TARATATATA CARTCARGAT TCTTGTCTCC ARGARACARA CCCARAGCCA ATRARAGCTA	1380
TAATGAACTT GGTTACAGGT GITACTTCTC TGACCTTCAA TCCTACTACA GAAATCTTGG	1440
CAATTGCTTC AGAAAAAATG AAAGAAGCAG TCAGATTGGT TCATCTTCCT TCCTGTACAG	1500
TATTITCAAA CITCCCAGTC ATTAAAAATA AGAATATITC TCATGTTCAT ACCATGGATT	. 1560
TITCTCCGAG AAGTGGATAC TITGCCTTGG GGAATGAAAA GGGCAAGGCC CTGATGTATA	1620
GGTTGCACCA TTACTCAGAC TTCTAAAGAG ACTATTTGAA GTCCAGTTGA GTCACAAGAG	1680
AAGCCTGTCT TGATATATCA TCTCAGAAAC TTTCCTGAAT ATGTGATAAT ATATGGAAAA	1740
TGATTTATAG ATCCAGCTGT GCTTAAGAGC CAGTAATGTC TTAATAAACA TGTGGCAGCT	1800
тттстттсаа ааааааааа аааааааааа аааааааа	1848

(2) INFORMATION FOR SEQ ID NO: 189:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1146 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

30 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 189:

AAAAAAACC CAGGGAACN TTGGGGGCCG CTTTNNNTTC CCCCTCCAGG CCATTGGGGA 60 ATTCTTCAAG TTAATCCTGC TTTGCTCTTG GCCAACAGGG CTTGTAGGGG GGAGAGACCC 120 AGGATCATCA AGGGGTTCGA GTGCAAGCCT CACTCCCAGC CCTGGCAGGC AGCCCTGTTC 180 GAGAAGACGC GOCTACTCTG TGGGGCGACG CTCATCGCCC CCAGATGGCT CCTGACAGCA 240 GCCCACTGCC TCAAGCCCCG CTACATAGTT CACCTGGGGC AGCACAACCT CCAGAAGGAG 300 GAGGGCTGTG AGCAGACCCG GACAGCCACT GAGTCCTTCC CCCACCCCGG CTTCAACAAC AGCCTCCCCA ACAAAGACCA CCGCAATGAC ATCATGCTGG TGAAGATGGC ATCGCCAGTC 420 TCCATCACCT GGGCTGTGCG-ACCCCTCACC CTCTCCTCAC GCTGTGTCAC TGCTGGCACC 480 AGCTGYCTCA TTTCCGGCTG GGGCAGMACG TCCAGCCCCC AGTTACGCCT GCCTCACACC 540 TTGSGATGCG CCAACATCAC CATCATTGAG CACCAGAAGT GTGAGAACGC CTACCCCGGC 600 AACATCACAG ACACCATGGT GTGTGCCAGC GTGCAGGAAG GGGGCAAGGA CTCCTGCCAG 660 GGTGACTCCG GGGCCCTCT GGTCTGTAAC CAGTCTCTTC AAGGCATTAT CTCCTGGGGC 720 CAGGATCCGT GTGCGATCAC CCGAAAGCCT GGTGTCTACA CGAAAGTCTG CAAATATGTG 780 GACTGGATCC AGGAGACGAT GAAGAACAAT TAGACTGGAC CCACCCACCA CAGCCCATCA 840

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	CCCTCCATTT CCACTTGGTG TTTGGTTCCT GTTCACTCTG TTAATAAGAA ACCCTA	AGCC 900
_	AAGACCCTCT ACGAACATTC TTTGGGCCTC CTGGACTACA GGAGATGCTG TCACTT	PAATA 960
5	ATCAACCTGG GGTTCGAAAT CAGTGAGACC TGGATTCAAA TTCTGCCTTG AAATAT	TGTG 1020
	ACTOTOGGAA TGACAACACC TGGTTTGTTC TCTGTTGTAT CCCCAGCCCC AAAGAC	CAGCT 1080
10	CCTGGCCATA TATCAAGGIT TCAATAAATA TTTGCTAAAT GAAAAARAAA AAAAA	1140
	ACTCGA	1146

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(2) INFORMATION FOR SEQ ID NO: 190:

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 906 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 190:

ACTOCCTOAC COAGGTOCCA GOCCTGGGAA COACCTACOG TGAGCCCTTT TGCAGATATA 60 GACTCATTTC ATCCTCAGAT GGTCCTTCAA GGTAGGTACT TTAGTCCCAT TTTAGAGATG 120 AGACGATTGA GGCCAGAGGG GTGNNGTAAC TTGCCTGGGG GCTCACGAGC ACAAAAGGAG CCGAGGCAGG ATCTGACCCT TGTTCTCTGG CCTCACTGCC CTCACTTTGC CATGACCCGA 240 AGITATGICC CTACAAAGCA ATGCATGGIC CAAGGYTCIT TITATIGIAT TITTATITIT 300 AAGGGTCCTG TTCAAAACTG GTGTGAGCTC TGAGGAGTCC TGAACCCTGG GTGCAGCATC 360 CTAGCATCCT GGGAGTCCTT TTCTGCCCAC ACTGAGCTGG GCTCCTCGAG GGGTGGGGCT 420 GCTGTCCCTG GAAGCCTGGC AGCAGCACTG TATCGGGTTG GCTGAAGCTG ARCGCCGTGG 480 GGTGCAGGGC TCCMGGAATC CCCGTTTGGC TGAAGGGGTT CCCTGTAGCC MGGGATGTTT 540 ATGAGGTCTC TCTGATGCCC CAGGCGCAGG ACATGTGTGC GGGTGGAGAA AAGCAGGCCC 600 TTTCAGTGCC AGCTCCACTC AATTTCTATG TGGACCAAGA ACGATAAACT TAAAAAATTT 660 TITTTCCTAA GGTATCTICA GAATATGGTG TATTTTTATG TGGAAAAGAA AAGTTATGAA 720 GGCAGCTGTT ACTITAAGAG AAAATTCATT AAAAGTCCTC GAGGTATGAA GATGACGGCG 780 TOCTTCTCAA TCATTTTGGC ATAACTTGAT TGTGGCTGTA ATTTTTTTTT TTTTTTTTGT 840 900 906 ACTCGA

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(2) INFORMATION FOR SEQ ID NO: 191:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1941 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 191:

10	(, 2020)	
	CTTCAGCTGA AGCCCAGGGA CCCCTTTTCC ACCCTGGGCC CCAATGCCGT CCTTTCCCCG	60
15	CAGAGACTOG TCTTGGAAAC CCTCAGCAAA CTCAGCATCC AGGACAACAA TGTGGACCTG	120
15	ATTCTGGCCA CACCCCCTT CAGCCGCCTG GAGAAGTTGT ATAGCACTAT GGTGCGCTTC	180
	CTCAGTGACC GAAAGAACCC GGTGTGCCGG AGATGGCTGT GGTACTGCTG GCCAACCTGG	240
20	CTCAGGGGGA CAGCCTGGCA GCTCGTGCCA TTGCAGTGCA GAAGGGCAGT ATCGGCAACC	300
	TOCTOGGETT CCTAGAGGAC AGCETTGCCG CCACACAGTT CCAGCAGAGC CAGGECAGCC	360
25	TCCTCCACAT GCAGAACCCA CCCTTTGAGC CAAYTAGTGT GGACATGATG CGGCGGGCTG	420
23	CCCGCGCGCT GCTTGCCTTG GCCAAGGTGG ACGAGAACCA CTCAGAGTTT ACTCTGTACG	480
	AATCACGGCT GTTGGACATC TCGGTATCAC CGTTGATGAA CTCAKTGGTT TCACAAGTCA	540
30	TTTGTGATGT ACTGTTTTTG NATTGGCCAG TCATGACAGC CGTGGGACAC CTCCCCCCC	600
	COTOTOTOTO TOCOTOTOTO GAGAACTTAG AAACTGACTG TTGCCCCTTTA TTTATGCAAA	660
35	ACCACCTCAG AATCCAGTTT ACCCTGTGCT GTCCAGCTTC TCCCTTGGGA AAAAGTCTCT	720
23	CCTGTTTCTC TCTCCTCCTT CCACCTCCCC TCCCTCCATC ACCTCACGCC TTTCTGTTCC	780
	TIGTOCTCAC CITACTCCCC TCAGGACCCT ACCCCACCCT CITIGAAAAG ACAAAGCTCT	840
40	GCCTACATAG AAGACTTTTT TTATTTTAAC CAAAGTTACT GTTGTTTACA GTGAGTTTGG	900
	GGAAAAAAA TAAAATAAAA ATGGCTTTCC CAGTCCTTGC ATCAACGGGA TGCCACATTT	960
45	CATAACTGTT TTTAATGGTA AAAAAAAAA AAAAAAATAC AAAAAAAAAT TCTGAAGGAC	1020
45	AAAAAAGGIG ACTGCIGAAC TGTGTGTGGT TTATTGTTGT ACATTCACAA TCTTGCAGGA	1080
	GCCAAGAAGT TOGCAGTTGT GAACAGACCC TGTTCACTGG AGAGGCCTGT GCAGTAGAGT	1140
50	GTAGACCCTT TCATGTACTG TACTGTACAC CTGATACTGT AAACATACTG TAATAATAAT	1200
	GTCTCACATG GAAACAGAAA ACGCTGGGTC AGCAGCAAGC TGTAGTTTTT AAAAATGTTT	1260
	TTAGTTAAAC GTTGAGGAGA AAAAAAAAAA AGGCTTTTCC CCCAAAGTAT CATGTGTGAA	1320
55	CCTACAACAC CCTGACCTCT TTCTCTCCTC CTTGATTGTA TGAATAACCC TGAGATCACC	1380
	TCTTAGAACT GGTTTTAACC TTTAGCTGCA GCGNCTACGT CNAWCGNTGT GTATATATAT	1440
60	GACGTKGTAC ATTGCACATA CCCTTGGATC CCCACAGTTK GGTCCTCCTC CCAGCTACCC	1500

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	CTTTATAGTA TGACGAGTTA ACAAGTTGGT GACCTGCACA AAGCGAGACA CAGCTATTTA	1560
_	ATCTCTTGCC CAGATATCGC CCCTCTTGGT GCGATGCTGT ACAGGTCTCT GTAAAAAGTC	1620
5	CTTGCTGTCT CAGCAGCCAA TCAACTTATA GTTTATTTTT TTCTGGGTTT TTGTTTTGTT	1680
	TTGTTTTCTT TCTAATCGAG GTGTGAAAAA GTTCTAGGTT CAGTTGAAGT TCTGATGAAG	1740
10	AAACACAATT GAGATTTTTT CAGTGATAAA ATCTGCATAT TIGTATTTCA ACAATGTAGC	1800
	TAAAACTTGA TGTAAATTCC TCCTTTTTTT CCTTTTTTGG CTTAATGAAT ATCATTTATT	1860
	CAGTATGAAA TCTTTATACT ATATGTTCCA CGTGTTAAGA ATAAATGTAC ATTAAATCTT	1920
15	GGTAAGACTT TAAAAAAAAA A	1941

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(2) INFORMATION FOR SEQ ID NO: 192:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2118 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 192:

AAATAATAAT AANAATAAAT AAAAATWAAG TGCTTAKTGT AACTCAGCGG ACAGGGCTCC 60 CAGCTGCTCT GGCACGTGGG ACACCYTCCA CCCTGCACAC AACAGGCATG CAAAGAGGAC 120 TOGATATOGT GOGGTAGAGT GCTTCTGGTG TGTTCACTTT AAGAAAACAT CTGCCAAGAG 180 AGAAGAGTGC CCAGGAAAGA CCAGGAAAAT ACAAGTACAT GGCTGCTTCA TACCATATAC 240 CCCAATTCTT TAAAGCAGCA AAAGGCACTT TTTTTTTCAG GCCAGAGTGA ATCTAAAACA 300 40 AACCTGGCTT TGCTTACAGG GAAGCTGTCC CAGAAGGACT GAGTGATGCC TCTTGTTCCC TAAGGTCTGG AGAGTCTTTG CAAGTTTCCA ACGACATTTC CAACCAGGTG GGAGAGACCA 420 GCAGTTGACG AGACAAGTCA GACCCAAAAA ACGACGCCAA GGTAGTGAGT GGGTGCCTAT 480 45 TTGGGAGTAG GATGATTTGA GGAAAACAGG AAGAAAAACC GGTCAGAAAG TGGCACTTTG 540 600 GAAGTGGAAA GCTGTTTGCA AATAGCAACT CTGGCTAAAG CGAAAATGTT AATCAAGTAG 50 AAAGTAAAAT TCAGGATCTT AGAAGCTCAT CCTTCTGATG AGAACTATTT TTTTTTCCGT 660 GAAGGAACTA TTATTACTTT AAAAGTGAGG GTAATTTACA TATGGGGTGT ATATATTCTA 720 AAAATAGTAA TAAAAGTACC TTTTATAAGC AATGTTGTGT GGCTTGTAGA AGAAAGCAGG 780 55 GAGGAAAAAA AGGCAGGCAA AACTAGTCTA GGTCTAGGCC CTAAAAATGA GCTTCCTTCC 840 CACTIGACIG GAAACGCCCA TGTGATTICT AGGCTGAAAA TAGGTAGGAT TTAACGAGTA 900

445

	CACCACCARGA TARGACTAC	960
	ACCTAGTTCC CTTCTGTCTC TGATTTCTGA TCAGCTGATG GAGCTGCTAG TAAGAGGGGC	-
	CGATCATGCT CCCAGACGAG TCCTTTGGCC TCTTGCTCTC CATCCCAAGC CTGACTCCTT	1020
5	CAGCAGCAGC CCCCTCCTTC TGTGTCCATC TGATGCAGGC AAGCAGGAGC AGTAAGAGGG	1080
	CATCCCATGT TCCAGTTCAC CTTCTATGGG GTGACTARGA GGTTCCCGGT AACTAGGGCA	1140
	GCCCARGCCC AGCAGGTTGC AAAAGCAGCT GCAAGCTTCA GAAACCCACT TCCTCCAACA	1200
10	CCAGGGAGGT GGCAGAGAGC CCATCCAAAA GCCCACTGGG AGAGGCATAA GATTCTGTGC	1260
	CAGGCCCCCA GGTCCCCTCT GTGTCAGGTA GGCTCTGCTA CTGGCCTCTG AAGTAAAGGC	1320
15	AAANACAAAC GGGCAGGGCA GGGTGGCAGG AATAAAAAAC TCTGGACAGA AACCCTTTTA	1380
	ATAAAGGAAA TTCCACCCCT CCCAATCCTT CCATGGAAGG GTGAGACCTT AATGTGATGT	1440
	AAGAGGAAGG TCTTCTCTGG CTTTCAGGGA AACAGCTGCA GCTGAAACTT AGGGGCCCAT	1500
20	TCCAGGGCAC TTTTCACCAC AGCCAGTGCA GCCGCTCCAA GTGCCACTGT CAGCCCCATC	1560
	ACTGCCAATT TCACAAAGCG GTTGGTCCTT GGCTTGGTCA GGACATCTTT TGTTCGATCT	1620
25	TCAGGCCGCA GAAGTCCCCG AANACCGCTG CCGCAGCACC ATATCAGGCC TCTGCTGGGC	1680
	TGATGCCAGC TCAAAGTCTT TGAAAGTAGA GGCTGCCGTC CTCTCAGCTT GCTGTTGGGC	1740
	AGCGGCCTCC CGAGCAAGTT CGGATGGGGG AAACTGAACA AAAAGGTCTC CTSTCTGCTG	1800
30	ATCAGTGTCT CATAGGGCAA GTCCTGAGGG ATCTGGGACA ACAGGTGGTG GACCGAGGCC	1860
	ATGTCACAGT CACAGTCCAG GACTTCCTGC TCGCGATACA ACACAATCAC GGCTGCAAAG	1920
35	TARATCGGCA TCAGTGGGTG GCAGGCCAGG AAGAAGTCAT ATAACCGCAC GACGTGCCTG	1980
	AAGTCAGACA GGACATGCCC AAACCAGGTG ATGAGCCAGC TGAGGGCAAA GATGGTCCCT	2040
	ACCTCAGCAC TCTGCATGAA GTCATGGAGC TCTGGATTCA CCTGGTCAAT GATGGGCATC	2100
40	AGATAGTTTA ATATATGC	2118

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(2) INFORMATION FOR SEQ ID NO: 193:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1538 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 193:

55
CCCGGTTCGG CTCTGTGTCA GCAGCCGGGC GGCGCTCGGG CGGGACATGG CAGCCTGTAC 60
ACCCCGGCGG CCTGGCCGTG GGCAGCCGCT GGTGGTCCCG GTCGCTGACT GNGGCCCGGT 120
60 GGCCAAGGCC GCTCTGTGCG CGGCCGNAGC TGGAGCCTTC TCGCCAGCGT CGACCACGAC 180

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	GACGCGGAGG CACCTCTCGT CCCGAAACCG ACCAGAGGGC AAAGTGTTGG AGACAGTTGG	240
5	TGTGTTTGAG GTGCCAAAAC AGAATGGAAA ATATGAGACC GGGCAGCTTT TCCTTCATAG	300
	CATTITIEGC TACCGAGGIG TCGTCCTGTT TCCCTGGCAG GCCAGACTGT RTGACCGGGA	360
	TGTGGCTTCT GCAGCTCCAG AAAAAGCAGA GAACCCTGCT GGCCATGGCT CCAAGGAGGT	420
10	GANAGCCANA ACTUACAUTT ACTATUAGGT GUTGATTGAT GUTCGTGAUT GUCCACATAT	480
	ATCTCAGAGA TCTCAGACAG AAGCTGTGAC CTTCTTGGCT AACCATGATG ACAGTCGGGC	540
	CCTCTATGCC ATCCCAGGCT TGGACTATGT CAGCCATGAA GACATCCTCC CCTACACCTC	600
15	CACTGATCAG GTTCCCATCC AACATGAACT CTTTGAAAGA TTTCTTCTGT ATGACCAGAC	660
	AAAAGCACCT CCTTTTGTGG CTCGGGAGAC GCTAAGGGCC TGGCAAGAGA AGAATCACCC	720
20	CTGGCTGGAG CTCTCCGATG TTCATCGGGA AACAACTGAG AACATACGTG TCACTGTCAT	780
	CCCCTTCTAC ATGGGCATGA GGGAAGCCCA GAATTCCCAC GTGTACTGGT GGCGCTACTG	840
	TATCCGITTG GAGAACCTTG ACAGTGATGT GGTACAGCTC CGGGAGCGGC ACTGGAGGAT	900
25	ATTCAGTCTC TCTGGCACCT TGGAGACAGT GCGAGGCCGA GGGGTAGTGG GCAGGGAACC	960
	AGTGTTATCC AAGGAGCAGC CTGCGTTCCA GTATAGCAGC CACGTCTCGC TGCAGGCTTC	1020
30	CAGTGGGCAC ATGTGGGGCA CGTTCCGCTT TGAAAGACCT GATGGCTCCC ACTTTGATGT	1080
	TCGGATTCCT CCCTTCTCCC TCGAAAGCAA TAAAGATGAG AAGACACCAC CCTCAGGCCT	1140
	TCACTGGTAG GCCAGCTGAG GCCCCAAGTG CCCAGGCTTG GTCACCGGGA AGAACAACTC	1200
35	TCATCCCACA ATTGCTGCAG AACTCTTCTC TCCCCATCAT GGGCCACAGT GGGTCTCTTA	1260
	ATTIGATIGT GOGGTTCTIT TIGTGGGGAG GGGTGGTATA ACTITTCTTC AGAAGACCCA	1320
40		1380
	CCTCTCCACC AAGGAACTGT GTTCAGCTGC CACAGGCCTG GAGGAGTTTC CTGGCCTGTC	1440
4	ACGTGAGGTT TGATCAGTAA ACCAGTGCAS GYTTGGCCAA AAAAAAAAAA AAAAAAAAAA	1500
45	ARRARARA ARRARARA ARAAAAAAAA AAACTCGA	153

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(2) INFORMATION FOR SEQ ID NO: 194:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1098 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 194:

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	AGACCCTGTC TCAAATAATA ATAATAATAA TAATCTTATT TTGGAGAATA AAGAGACCTS	60
	TOGATTIGAG GIGCCATTIG GGTAGAAAGA AAAGACGITT ACACCGAGAA ATAGICTGIG	120
5	TIGCCCTGAA GGAGCAGAGG GATGCATCGC TGGAGGTGAC CTACAGTTGA AGAAGACTCA	180
	TTATGACAGA CCTTGTCCTT CTTCCTTGTG GAAAGTGTTT CCTCTGCTGC TACTGCTCAT	240
	GAGACTOTTO COCCTCCCTG TCCCAGGGAA CCAAAGGGCT TINCTACCAC ACCOTTTCTT	300
10	NGCCCCCCCC CTCCCATGTC TGCTGTGCCT TTGTACTCAG CAATTCTTNG TTTGCTCCCA	360
	TTATCTTCCA GCCGGATACA GAGTGAATAG TTAACCACAC TTAGGTCAAA TAGGATCTAA	420
15	ATTITIGITC CIGCICCNGT GTAAAGAGGC CAGIGITIGI GIGTIGCAAG CAGCCTIGGA	480
	ATAGTAACTC TTCTCATTTG TTTGGGATCT GGCCAMCAAG TTCCAGAATG ATACACGGAT	540
	CAGTGCAGAA GTTCATCAGG CTCTCGGACC TTAGGGCTGT TGGAGAAGGC TTCAGCAGCA	600
20	GAACTGATGG TKAWKGYTCG TGTTCTCCAT CCTCAACTTT CTTTGCTTCG ATCATACACA	660
	AGAATACATT TOGAAGGGCA AAAAATGAAC ACTGTTGTTC ATTGCAGCCG TGTTTTGTGA	720
25	CACAGATGCA CAGTCTGCTG TGAAGACCTT CTCTCAAGTG GSATYTGGGA GTCCATGCCA	780
	GATCATGGTG CTTCATGAGA GACTGACAGC TATCAGGGGT TGTGGCACTT AGTGAGGACT	840
	CTCCTCCCCC AGTGTGTGCT GATGACACAT ACACACCTGA CAATAGCTTG AGTCTTCTCT	900
30	GTTCCTTTTA CTCTGTAGCC AACATACACA TGATTTAAAA CCCTTTCTAA ATATCTATCA	960
	TOGTTCATCC TTGTCCAAAT GCAGAGTCAG AGCTATTTGT ACTTCATTAT TATTTCCAAG	1020
35	GCGAATAGTT GGCTTTCTTT TTGCAAAAAT AATTAAAGTT TTTGTATGTT GCAAAAAAAA	1080
	AAAAAAAAA CTACGTAG	1098
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40	101 THE TOTAL POR STO TR NO. 195.	
٠.	(2) INFORMATION FOR SEQ ID NO: 195:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1001 base pairs	
43	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 195:	
	GAATTCGCCA CGAGATAGCT TGCATCTCAT CCCAGTAAAA CCACTTATTT ATAACATATC	60
	AACGTATIGA CAAGGTIGAA GAGCAAGATT GITCTGAGGT GAGATGCAAA TITCAAAGGG	120

GTGAGCACTA ATTGTTCCAG TGATTGTTTA TTTATTGGCT AGGACATAAT TACTCTCTTT

GAGGTTACAC ATCTGCCTCC AGGTTCCTGT GTGCTTGTGC CCTTGGGATC AGGCCAGGGC

AGACTGTGAT CACTGAGATT CAAACTCCCA GARTAATCAG CAAGAGCTTT CTAGAGACCA

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•	AGGCCAGGCC TGATCCCTGA GGGATGCATG AGAAGGCTTG GAATCTCATT CTGCTATGGT	360
_	GGCTCTCTCT TGATCTTCTT GGAGTAGCAA AAACAGCAAT GTGGGCCCCAA TGGTGTGGCC	420
5	TAAATGATCA CAAAGGTAAA TGAGTAAAGG GCTCAGCAGA TGAGTAAGGA GCCTTGTCCT	480
	GAGAAATTAG CACTGGGCTC TGCATTCAGA AACATGTGAT AAGCATTGCC CATTGCACAT	540
10	TGCCTTTATT GTGTAAGGAC ATGAAATTCC AGTTTTGCAT AGCTAGTGAT GAATACCTGA	600
	AGGGAATTGC AGACATATTT TATTTTATTT TTAATTGACA GATGGAATTG TATATATTTA	660
15	TCATGTACAT AATCATGCTT TAAAATATGT ACATTATGGA ATGGCTAAAT CAAACTAACC	720
13	TAGGCATTAT CTCATATAAT TGTCATTTTT GTGGCGAGAA GACTAAAAAT CTACCCTTTC	780
	AGCATTITTA AAGAATACAA TGTGTTTTAT TAACAACAGT CACCATTTGG TACACTAGAT	840
20	CTCTTGAACT TCTTCCTCTT ATCTAACTGA GATCTTGTAA CCTTTGATAA CAGCTCCCAA	900
	GCCCTTCCCC AACCACTGCT CCACCCGTGG TAACCACCAT TCTATTCTCA ACTTCCTGGT	960
25	AATCACCATT CTAGACACAG GGAAGACTCT CTACCCTCTG A	1001
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(2) INFORMATION FOR SEQ ID NO: 196:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1443 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 196:

60	T GTCATAAGAA	TAAATTATTT	ATATTATTTT	САААТАТААА	ATAGGTCATG	ATAAACTGAA
120	T TCTKTGGAAG	TGGTTAGCAT	GGAAAAAATG	CTTTAATAAT	CCATATTTTG	ACGATOGTGG
180	T GTGGAAATGT	ACCTGCATAT	ATTTATTTCT	ATTTTCTAGG	GATAGTAGAC	GTGGTCATCA
240	G TGCTAATGGT	CATCAAAATG	AACTCAGAGG	AATGGCAGCT	AGATTTATWT	GTACTACTTI
300	C ARGGGCAGGG	TCAATCAAGC	TIGTARGCCT	CICTYCICTT	CTTTGTCTTG	GTAATATGGC
360	C CGTCTCCACT	CCCCTGACCC	CCAGCGTCTG	TTGSCAGACG	AACTTGTCCT	CCGTACAGTO
420	IC TECCTECCIT	ATTCACCTTC	GCYTACCCTG	GCCCCTTGAT	TGGAGGAGGA	CTCTGTGTCC
480	TA CACCATTGAT	TCCTTGCTTA	GGATCTGAAA	GTGCAATAAC	r gggaagagcc	GTACTGAACT
540	AA AGATCTCCTT	ATGTTATGA	: ACCACCATGO	CGTGGGCAAC	A CTAGCATTAC	CTAGGAGACI
600	CA CATCAGCACT	TGCATCAGC	AGTAGAAATO	G AGTGCAAGGG	A CCTACCGGT	CCAGAAACC
660	CT CCCAGCTGTT	GATGTCATC	TGACCAAGT	TOGGGGAAAA	A AGTAAACCTY	TGGGGATCT

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	TOTANGAGOC CAGATGTCCA GAGTATTGTC TCACCTTGAT CCCTCAGGCC AGAAGACCTG	720
	TGAAAAAGCC ACACTGGTTC AGGGACTCAC TGGACGGTTT TGTGTCCAUT YTAACTTTGCA	. 780
5	COGTOTOTAC COCAGAGTOG ACTORRATCO TOARGTORIO CTOTGRARDAT TORRESTORGA	840
	AATTATAAAA GGGCTTTGGC AATATGTTAG CCCAAGAATT TGGCTTCTTC CAGAAATTGT	900
	GCCGACNITA ACAGTGCCTT AAATGATGGT AAAACTITITA AGATTICTIAA AAGGTGGCA	960
10	TIGGAGATAC GITGACTITT ATTAAACMAC CTATAGTTST TTAATGATTT CTAAAAAAAT	1020
	ATCTGGAGCT CAGGGGTTCA ACTGAGGGAA CACATGTTGA GRATCATTGT TTALTAATTA	1080
15	ANTIGOCAGGT AACCCGTTGA AATTATCAAA AACATCTTCC ACGTACCAGA AAGCACCTCA	1140
	GAGGATAGTT CTGTTATGGA GAAGATGAAA TGGTTIAGTA GTGTAGGAAC TATGGAAAGG	1200
20	TGAGCTTAGA TTTGGATAGT AAAACCTCAA GACCCTATTT AAAAAGTATT TTATGAATGC	1260
20	AGCATAAATA ATTTAATTCA GTGTTAANAT GCCAASGCTA GTATATTGAG CTGAATGTGA	1320
	AAAGAAACTC ACATTGOGAG AATGCCACCT TITCCITATA AGATAGCTIT GAAGATACCA	1380
25	TITTAGACAG ATGGAAATTG AATAGCTITA GAAAAGGCAA ATGTITGATC TIGGGGAAAA	1440
	AAA	1443

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(2) INFORMATION FOR SEQ ID NO: 197:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1282 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: dcuble

(D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ☐ NO: 197:

GAAAAAAAA AGTATGACCC AGTAGCTAGG CACCTGTGGC CCCGCCAAGT TGACACATAA 60 AATTAACTGT CACAGTATCA TCTTAGAAGT GAAAGAAGCC CCTTTATCCT GCAGTGCCCC TCTACCACCA CCTACTGACA AAGAACATGG TGCTATCTGG CATGGGAGAA ATGTTCAGTT 180 TGCTATGGCT TGTATGTGTC CCCTCAAATT CAAGTGTTGC CAATGTGACA GCATCAAGAG GTGGGGTCTT TAAGAGATCA CTAGGCCATG AGGGATTCTC TTAGGACTGG GATGAAGGCC 300 CATAATAAAA GAGGITTCAG GGAGCATCCT GCTAGCTTGC CTTCTGTATG TGAGAACACA 360 GCAAGAAGC CCTAGTCAAC AAGTGCCAGC TCCTTGATCT TAGACTTCCC ATCCTCCAGA 420 ACTGTGAGAA ATACATTTCT GTTCCTTACA AATTACCCAG TCTCCTGTAT TCTGTTATAG 480 CAGCACAAAA TGAAGATACC ATACCTGAAC ACCTGAACAT TCTTCACAAG GTAGTAAATG 540 CACTGCTTTA TTCTGGTCTC AGTATTGTGT GCTTAATAAG GAAATGAGAA AGGSTGGATC 600

•	AGGGCATAGG ATGAACAAGT TACTGCTAGA CCTCTCACAA TGCCACTAAT GGATAAGATT	660
_	GTATTTCAT CATTNCTTGT CTCTTCGGAA GCTAACACCA TGCTATAATA GGCALTAAAT	720
5	AGATGTCTAA AAACACCTTA AGTATTTGTC TAGAAATTTG GTGCATTGTC CAGAAAGAAC	780
	CAAAATTCMA AATAATTTCA AAGGGCCTAA AGCACTAKTT AATCIGAAATT CATTAGTFFF	840
10	TARTOGTACT ACCACTCTCA ARTITAAAAT GTCATCTTAC GTTCCTCTTC CTCGCATTGG	900
	ATTITATIGCT AAAACCIGGT AAACACTITA ATCCYTTICA ATTCCATTAC CACTGCTCTT	960
15	GTCCAGAATT ACTOGCAGAC TAATAGTCAC CTGACTTCTC CCCCTGCATC CCGATTTGCT	1020
13	GTCTAATTCT GGTTACAAAT AAGTAACTGC CAAACTAATC TTTCTAAAAA GCAAGACTGA	1080
	TOTOGTCACT COTTTGCTCA ACAATGTAAA AGCTCCCATT GTCTCCCAAA TAAAACCAGC	1140
20	TTTCCACTGT GTATACAATA CATCCATGAT CTGTATCCAG CATCATTTTG TATTICCTCA	1200
	CTTTATACAC CACCCCCCAT GCCACATCAA ATTAAATTAT CCTGATAAAT GCAACTGCAA	1260
25	AAAAAAAAA AAAAAAACTC GA	1282

(2) INFORMATION FOR SEQ ID NO: 198:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 951 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double

(D) TOPOLOGY: linear 35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 198:

40	ATTTCGGAAC GAGGACTGAA GTGGGAGCGG CGGCAGGGTA GAAGALAGAA GGGGGATCTA	60
40	TGTGGTAACT AAAGAATGTT TCTGTTTTGT TAATTATTGT GTGTGTGTGG TTTTATTGTT	120
-	TGCTTAAGAG AATCAAAAAC TGAAAAAAAT GAGAATACAG GAAATGGCTC TTGTTTATTT	180
45	TITITGCTGTG THIACACCTT GTTAATGCTC TACTGTCTTT GTTTCAAGAG AGATTTGTTC	240
	ACTOCCCAGO TOGITTIGIG TOCTGAGCCC TATGCCCAGO CCACCITATA AATCATGCCT	300
50	GITTAGATGT TIGATTTIGT TCTGTTTGCT ATTGTTATCT TAAAGGTGTA TAACCCTGAC	360
50	ATGCCAGACA TCAAATTAAG CTCAAATTAA GCTCTCGTTT AAATGTTAA ACACCTAATT	420
	TATATTCTAA TIGATCCCAG CCACTGATGC ATGTACTTTA GCTACTTCTG CTAAATAAGC	480
55	ATATTAATIT TOCACATCAG GCCATCAGAT CITGAGAACC AACAGITAIC CAGAATTCCG	540
	TGTCTACTAA TGTTTCACCT GCATGCAGCC TTCATTAATT TTGTAGCAAA ATATAAAGTG	600
60	ATCATTATGT AGTITICIGGA TTAAAAAAAT TIGTGTGTGA AGTIGCTITG CAAAGTGCAT	660

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•	GTGGAATTAA TGGGACAGTG TGCCCTTTGT GTTAGATGTT AGAGCAAAAG AAAGGGCTTA	720
	TAGTGTTAGT ATTGGAGCAC TTTGAAGATA GATATTTTCA GAAAAGATGT AGGATTTAAA	780
5	AGITAAATTT TAAATTTTAG AAAAAGATAT GATGGCAATT GGAAATAGTC ACAATGAAGT	840
	TCTTCATCCA GTAGGIGTTT AACAGTGTTA TTTTGCCACT GGTAATGTGT AAACTGTGAG	900
10	TGATTTACAA TAAATGATTA TGAATTCAAA AAAAAAAAAA	951
15	(2) INFORMATION FOR SEQ ID NO: 199: (i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1740 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 199:	
25	TTATTATAAT AATGATGATG ATTCCAAGGA AAAAACCTAC AGCGAATGTT CCATTTCTAC	60
23	CCCGCACGCA GACACTCTCC CTAACACTGA TAACCTGAGC CCCCAGCACT GGACGGAAGA	120
	ATGCTGGCGT CTCCGTGTGT ACTGGTTCAG GGTTCTGGCC CCAGCCTTGT CAGGACCCCC	180
30	TOGTGTCCAG AGCCCCCACC CCTCCCGCAA CAAGCAGCTG ATGCCCCAGT GATTCTCTAT	240
	ACATTITICA CCTCGCCAA TATGTCCAGG AAAACTGCTT ACTTCTCTTT TCTTGCCTGG	300
25	AGCCTTCATT GTTCACCCTT ACGTTGCAAT ATAGGAATTA ATGCTACAAA ATAAAAGTAA	360
35	AGCTTACCTG AAAAGTGCAT AGTTTGGGGC AATGGTATCT ACATCTCCCA CTGTGGGAAA	420
	ACCAGCAAAG CATCAAAACT CTCAATTCTC CTGTTACCRA ATGCAGATCT GAATTATAAG	480
40	ATGITTATGT TIGACCATTG TTICAACAAT GGGATTITGT TACGAATTAT CCCTTTAACT	540
	GAAACCCTCA GTTTTACTGT TTACATTATT AGGAAAACAG GGATATCTTT TGAATCTAAA	600
	AATTIGATGT ACAGCATGTG ATTITTGAAG TITACATGTA AAGTCACAGT ATAGGTGAAA	660
45	TAACGTTIGT CATATTTIGA GACGTATCCT GCAGCCATGT TTTTACGTGA GTGTTTTAGT	720
	CAAAGTACAT GGTAGACAGT CTTTCACAAT AAAAGGAAAA GGATTTTTTT TCCTCCAAAT	780
50	GTACATTTAT CAACCTAATG ATTGATTTTT TTAAAAAGAG ATTTCGCCCC AGTCTGGTTT	840
	ATGAAAGITC ATTGCCCTAA ACTGTGCTGA TTGTTTTTAA TCAAGTTATA AATTTCCAAC	900
	CTAGATCATG TATCTACCAA CTCTCCTGCA TTTTCCAAAA GGCATTGAGC TTAAATATTA	960
55		1020

GTCTTGCTTA GAGTAGGTTA TCCACTTACA TGCTGCGCTA AAGCCATGCC TTTGAAACTC

CTTGTTTAAA ACATGATATG ATTTTTGTGG GCAGTTTCAG AAAAGAAAAC AAACAAACAA

AAATCGACCC TITAATTATT ACTIGCAACT CAACAGATCT CCCTGCCGTA CTGCCTTTTC

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	CAGGAACTIT ACTICAGGC TGTCCAGATT GCAGTTGTGC CCCGTGTATG TGGATCTAGT	1200
	TCACAGAGTC TTTGGAAGCC AGCAGTCGTG CCCTCCGTAT ACTGTCCACT CATTTTATGT	1260
5	AGATTTOGTA TCCTCAGCAG CCAGTGTTAA CACCACTGTC ACGTAGTTAN CAGATTCATC	1320
	TITTATGTAT TTAAAGTAAT CCATACTATG ATTIGGTTTT TCCCTGCACC ATTAATTCTG	1380
10	GCATCAGATC AGTTTTTGTG TTGTGAAGTT CTACTGTGGT TTGACCCAAG ACCACAACCA	1440
	TGAGACCCTG AAGTAAAGAT AAGGTACACA TACATTATTT GAGTAACTGT TTCCTTGGGG	1500
	GCCAATCTGT GTATGCTTTT AGAAGTTTAC AGAATGCTTT TATTTTTGTC TATAACAAAC	1560
15	AGTOTOTOAT TTATTTCTGT TGATAAACCA TTTGGACAGA GTGAGGACGT TTGCCCTGTT	1620
	ATCTCCTAGT GCTAACAATA CACTCCAGTC ATGAGCCGGG CTTTACAAAT AAAGCACTTT	1680
20	TGATGACTCA MAAAAAAAA AAAAAAAAMC YCGGGGGGGG GCCGGTAACC CATTTNNCCC	1740

25 (2) INFORMATION FOR SEQ ID NO: 200:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1707 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 200:

GCTTATAGAA GGGAGAGGAG CGAACATGGC AGCGCGTTGG CGGTTTTGGT GTGTCTCTGT 60 35 GACCATGGTG GTGGCGCTGC TCATCGTTTG CGACGTTCCC TCAGCCTCTG CCCAAAGAAA GAAGGAGATG GTGTTATCTG AAAAGGTTAG TCAGCTGATG GAATGGACTA ACAAAAGACC 180 40 TGTAATAAGA ATGAATGGAG ACAAGTTCCG TCGCCTTGTG AAAGCCCCCAC CGAGAAATTA 240 CTCCGTTATC GTCATGTTCA CTGCTCTCCA ACTGCATAGA CAGTGTGTCG TTTGCAAGCA 300 AGCTGATGAA GAATTCCAGA TCCTGGCAAA CTCCTGGCGA TACTCCAGTG CATTCACCAA 360 45 CAGGATATTT TTTGCCATGG TGGATTTTGA TGAAGGCTCT GATGTATTTC AGATGCTAAA 420 CATGAATTCA GCTCCAACTT TCATCAACTT TCCTGCAAAA GGGAAACCCA AACGGGGTGA 480 50 TACATATGAG TTACAGGTGC GGGGTTTTTC AGCTGAGCAG ATTGCCCGGT GGATCGCCGA 540 CAGAACTGAT GTCAATATTA GAGTGATTAG ACCCCCAAAT TATGCTGGTC CCCTTATGTT GGGATTGCTT TTGGCTGTTA TTGGTGGACT TGTGTATCTT CGAAGAGTAA TATGGAATTT 660 55 CICITTAATA AAACTGGATG GGCTTTTGCA GCTTTGTGTT TTGTGCTTGC TATGACATCT 720 GGTCAAATGT GGAACCATAT AAGAGGACCA CCATATGCCC ATAAGAATCC CCACACGGGA 780

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	CATGTGAATT ATATCCATGG AAGCAGTCAA GCCCAGTTTG TAGCTGAAAC ACACATTGTT	840
	CTICTGTTTA ATGGTGGAGT TACCTTAGGA ATGGTGCTTT TATGTGAAGC TGCTACCTCT	900
5	GACATOGATA TTOGAAAGCG AAAGATAATG TGTGTGGCTG GTATTGGACT TGTTGTATTA	960
	TICTICAGTI GGATGCTCTC TATTITTAGA TCTAAATATC ATGGCTACCC ATACAGCTTT	1020
	CTGATGAGTT AAAAAGGTCC CAGAGATATA TAGACACTGG AGTACTGGAA ATTGAAAAAC	1080
10	GAAAATCGTG TGTGTTTGAA AAGAAGAATG CAACTTGTAT ATTTTGTATT ACCTCTTTTT	1140
	TTCAAGTGAT TTAAATAGTT AATCATTTAA CCAAAGAAGA TGTGTAGTGC CTTAACAAGC	1200
15	AATCCTCTGT CAAAATCTGA GGTATTTGAA AATAATTATC CTCTTAACCT TCTCTTCCCA	1260
	GTGAACTTTA TGGAACATTT AATTTAGTAC AATTAAGTAT ATTATAAAAA TTGTAAAACT	1320
	ACTACTITGT TITAGITAGA ACAAAGCTCA AAACTACTIT AGITAACTIG GICATCTGAT	1380
20	TTTATATTGC CTTATCCAAA GATGGGGAAA GTAAGTCCTG ACCAGGTGTT CCCACATATG	1440
	CCTGTTACAG ATAACTACAT TAGGAATTCA TTCTTAGCTT CTTCATCTTT GTGTGGATGT	1500
25	GTATACTITA COCATCTITC CITTIGAGTA GAGAAATTAT GTGTGTCATG TGGTCTTCTG	1560
	AAAATGGAAC ACCATTCTTC AGAGCACACG TCTAGCCCTC AGCAAGACAG TTGTTTCTCC	1620
	TCCTCCTTGC ATATTTCCTA CTGAAATACA GTGCTGTCTA TGATTGTTTT TGTTTTGTTG	1680
30	TTTTTYGAG ATCACGYTAC TGGGCTC	1707
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	(2) INFORMATION FOR SEQ ID NO: 201:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 779 base pairs	
40	(B) TYPE: nucleic acid	
.,	(C) STRANDEINESS: double	
	(D) TOPOLOGY: linear	
	A CONTRACT DESCRIPTION SEC ID NO: 201:	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 201

	(XI) SEQUENCE PERSONNEL	
45	CTGTCCCCAG TGTTTCCAGG TAATGACTTG GCACTCCAGA GAAAGTTTCA TRCTGTTGCG	60
	TGTGGTGGCT CCAAGCCAAG CACCTGGCAT GCAGGTCAGC CCTTCCCAGC GGGCGTGGCG	120
50	TOGTOCTCTT CACAGATGCC ACGTTGCAGC CCCAAGGCCT CACCATTTTG CGTTTTTTAG	180
	ARACCCATTT TCTTGGTCAT TTATARAGCT GCTTTATAGA TATCTTTGAT CCTGGCATGC	240
	CTTGGTTTCC TCTCCCTTCC CTCTTTCCAA TCCTGGTTTC CTAACCTCCT CTTGTAGTAA	300
55	TTCTCAACTC AACTCAAAGT CCCAAGAATT TGGAATGGTA GGATGCTGTG CGGGGAGCTC	360
	GAGGCTGAGG CATAATCACT GCTTCGGTTC TGCTCATCAG GGGACACGCT CCCTTACTCA	420
60	TOGCAGCCAT GITTGATTGT CACAGAGCCC CCCGAATACT CTGTCTATAG TGACACACTG	480

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	TAGGTGTCAT AAATTTTAAG AAACCTGCTT TTAAGTACTA TTTATAGGTT TTTCTGTTAT	540
	ACTTGCAACC TAGTTTTAAA ATACATGAGG ATTTTATGAA AGCTTTATAC AGACATTTAT	600
5	AGGAAACTCA TTCTTTGATT TTAGGTGCCA TTTAAATTGA TAACACTTAC TTTATAAAAA	660
	GATGCTTTTT GTCTGGATAG AGCCTTATAG TTTAAAATAT CTTCATATAT TGCCATTTGA	. 720
10	тсаратарат ттсттастта сарарарара арарарара арарарара арарастсса	779
15	(2) INFORMATION FOR SEQ ID NO: 202:	
13	(2) INFORMATION FOR DDQ 22 HOT DDG	
	(i) SEQUENCE CHARACTERISTICS:	

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(A) LENGTH: 1617 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 202:

25	GGCACAGCTT TCTGTCTCTT CCTCGCTCCC TCTCTTTCTC TCCTCCCTC	60
	TGCATAAAGT CTCTGTCGCT CCCGGAACTT GTTGGCAATG CCTATTTTTT GGCTTTCCCC	120
	CGCGTTCTCT AAACTAACTA TTTAAAGGTC TGCGGTCGCA AATGGTTTGA CTAAACGTAG	180
30	GATGGGACTT AAGTTGAACG GCAGATATAT TTCACTGATC CTCGCGGTGC AAATAGCGTA	240
	TCTGGTGCAG GCCGTGAGAG CAGCGGGCAA GTGCGATGCG GTCTTCAAGG GCTTTTCGGA	300
35	CTGTTTGCTC AAGCTGGGCG ACACATGGCC AACTACCCGC AGCCTGGGAC GACAAGACGA	360
	ACATCAAGAC CGTGTGCACA TACTGGGAGG ATTTCCACAG CTGCACGGTC ACAGCCCTTA	420
	CCGATTCCCA CGAAGCCCC AAAGATATCT CCGATAAACT GAGAAAAGAA TCCAAAAACC	480
40	TCAACATCCA AGGCAGCTTA TTCGAACTCT GCGGCAGCGG CAACGGGGCG GCGGGGTCCC	540
	TOCTOCCOGC GTTCCCGGTG CTCCTGGTGT CTCTCTCGGC AGCTTTAGCG ACCTGGCTTT	600
45	CCTTCTGAGC GTGGGGCCAG CTCCCCCCGC GCGCCCACCC ACACTCACTC CATGCTCCCG	660
	GAAATCGAGA GGAAGATCCA TTAGTTCTTT GGGGACGTTG TGATTCTCTG TGATGCTGAA	720
	AACACTCATA TAGGATTGTG GGAAATCCTG ATTCTCTTTT TTATTTCGTT TGATTTCTTG	780
50	TGTTTTATTT GCCAAATGTT ACCAATCAGT GAGCAAGCAA GCACAGCCAA AATCGGACCT	840
	CAGCTITAGT CCGTCTTCAC ACACAAATAA GAAAACGGCA AACCCACCCC ATTTTTTAAT	900
55	TTTATTATTA TTAATTTTTT TTGTTGGCAA AAGAATCTCA GGAACGGCCC TGGGCACCTA	960
"	CTATATTAAT CATGCTAGTA ACATGAAAAA TGATGGGCTC CTCCTAATAG GAAGGCGAGG	1020
	ACACCAGARGA CYCAGGGGAA TGAATTCAAG AGAGATGTCC ACGGACGAAA CATACGGTGA	1080

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	ATAATTCACG CTCACGTCGT TCTTCCACAG TATCTTGTTT TGATCATTTC CACTGCACAT	1140
	TTCTCCTCAA GAAAAGCGAA AGGACAGACT GTTGGCTTTG TGTTTGGAGG ATAGGAGGGA	1200
5	GAGAGGGAAG GGGCTGAGGA AATCTCTGGG GTAAGAGTAA AGGCTTCCAG AAGACATGCT	1260
	GCTATGGTCA CTGAGGGGTT AGCTTTATCT GCTGTTGTTG ATGCATCCGT CCAAGTTCAC	1320
••	TECCTPTATT TTCCCTCCTC CCTCTTGTTT TAGCTGTTAC ACACACAGTA ATACCTGAAT	1380
10	ATCCAACGGT ATAGATCACA AGGGGGGGAT GTTAAATGTT AATCTAAAAT ATAGCTAAAA	1440
	AAAGATTTTG ACATAAAAGA GCCTTGATTT TAAAAAAAAA AGAGAGAGAG ATGTAATTTA	1500
15	AAAAGTTTAT TATAAATTAA ATTCAGCAAA AAAAGATTTG CTACAAAGTA TAGAGAAGTA	1560
	TAAAATAAAA GTTATTGTTT GAAAAAAAAA AAAAAAAA	1617

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(2) INFORMATION FOR SEQ ID NO: 203:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1974 base pairs(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 203:

GAATTCGGCA CGAGGCTGAG GGAGCTGCAG CGCAGCAGAG TATCTGACGG CGCCAGGTTG 60 CGTAGGTGCG GCACGAGGAG TTTTCCCGGC AGCGAGGAGG TCCTGAGCAG CATGGCCCGG AGGAGGGCT TCCCTGCCGC CGCGCTCTGG CTCTGGAGCA TCCTCCTGTG CCTGCTGGCA 180 CTGCGGGCGG AGGCCGGGCC GCCGCAGGAG GAGAGCCTGT ACCTATGGAT CGATGCTCAC 240 CAGGCAAGAG TACTCATAGG ATTTGAAGAA GATATCCTGA TTGTTTCAGA GGGGAAAATG 300 GCACCTTTTA CACATGATTT CAGAAAAGCG CAACAGAGAA TGCCAGCTAT TCCTGTCAAT ATCCATTCCA TGAATTTTAC CTGGCAAGCT GCAGGCCAGG CAGAATACTT CTATGAATTC 420 CTGTCCTTGC GCTCCCTGGA TAAAGGCATC ATGGCAGATC CAACCGTCAA TGTCCCTCTG 480 CTGGGAACAG TGCCTCACAA GGCATCAGTT GTTCAAGTTG GTTTCCCATG TCTTGGAAAA 540 CAGGATGGGG TGGCAGCATT TGAAGTGGAT GTGATTGTTA TGAATTCTGA AGGCAACACC 600 ATTCTCCAAA CACCTCAAAA TGCTATCTTC TTTAAAACAT GTCAACAAGC TGAGTGCCCA 660 GCCGCGTGCC GAAATGGAGG CTTTTGTAAT GAAAGACGCA TCTGCGAGTG TCCTGATGGG 720 TTCCACGGAC CTCACTGTGA GAAAGCCCTT TGTACCCCAC GATGTATGAA TGGTGGACTT 780 TGTGTGACTC CTGGTTTCTG CATCTGCCCA CCTGGATTCT ATGGAGTGAA CTGTGACAAA GCAAACTGCT CAACCACCTG CTTTAATGGA GGGACCTGTT TCTACCCTGG AAAATGTATT 900

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	TSCCCTCCAG GACTAGAGGG AGAGCAGTGT GAAATCAGCA AATGCCCACA ACCCTGTCGA	960
	AATGGAGGTA AATGCATTGG TAAAAGCAAA TGTAAGTKTT CCAAAGGTTA CCAGGGAGAC	1020
5	CTCTGTTCAA AGCCTGTCTG CGAGCCTGGC TGTGGTGCAC ATGGAACCTG CCATGAACCC	1080
	AACAAATGCC AATGTCAAGA AGGTTGGCAT GGAAGACACT GCAATAAAAG GTACGAAGCC	1140
10	AGCCTCATAC ATGCCCTGAG GCCAGCAGGC GCCCAGCTCA GGCAGCACAC GCCTTCACTT	1200
	AAAAAGGCCG AGGAGCGGCG GGATCCACCT GAATCCAATT ACATCTGGTG AACTCCGACA	1260
1.5	TCTGAAACGT TTTAAGTTAC ACCAAGTTCA TAGCCTTTGT TAACCTTTCA TGTGTTGAAT	1320
15	GTTCAAATAA TGTTCATTAC ACTTAAGAAT ACTGGCCTGA ATTTTATTAG CTTCATTATA	1380
	AATCACTGAG CTGATATITA CTCTTCCTTT TAAGTITTCT AAGTACGTCT GTAGCATGAT	1440
20	GGTATAGATT TICTTGTTTC AGTGCTTTGG GACAGATTTT ATATTATGTC AATTGATCAG	1500
	GTTAAAATTT TCAGTGTGTA GTTGGCAGAT ATTTTCAAAA TTACAATGCA TITATGGTGT	1560
25	CTGGGGGCAG GGGAACATCA GAAAGGTTAA ATTGGGCAAA AATGCGTAAG TCACAAGAAT	1620
23	TTGGATGGTG CAGITAATGT TGAAGTTACA GCATTTCAGA TTTTATTGTC AGATATTTAG	1680
	ATGTTTGTTA CATTTTTAAA AATTGCTCTT AATTTTTAAA CTCTCAATAC AATATATTT	1740
30	GACCTTACCA TTATTCCAGA GATTCAGTAT TAAAAAAAAA AAAATTACAC TGTGGTAGTG	1800
	GCATTTAAAC AATATAATAT ATTCTAAACA CAATGAAATA GGGAATATAA TGTATGAACT	1860
35	TTTTGCATTG GCTTGAAGCA ATATAATATA TTGTAAACAA AACACAGCTC TTACCTAATA	1920
<i>)</i> ,	AACATTITAT ACTOTITGTA TGTATAAAAT AAAGGTGCTG CTITAGTTTT CTGA	1974

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(2) INFORMATION FOR SEQ ID NO: 204:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1057 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 204:

50 COGCCTTCCG GGCAACCGT TCGTCCCAAC NCGGGAAAGG GTCCTGGAGN CGGGAACTAG 60 GAGCCTCGGA AGTCCAAGGG CGGAGCGCCC TTTGCTAATA AGCCAATCAG AACGTGAGAC 120 SCICCOSTICS GINCOSTICCG TOGASCICCG GGTGGASTCT GGGTGACTTG GCTGGCGGGA 180 55 TCAAGTGCAG CTGCTTCAGG CTGAGGTGGC AGATAGTGAG CGCTGGTGGC GGAGTTAAAG 240 TYAAAGCAGG AGAGTAATWA TGAATAGCGC AGCGGGATTC TCACACCTAG ACCGTCGCGA 300 60

457

	GCGGGTTCTC AAGTTAGGGG AGAGTTTCGA GAAGCAGCCG CGCTGCGCTT CCACACTGTG	360
•	CGCTATGACT TCAAACCTGC TTCTATTGAC ACTTCTTCTG AAGGATACCT TGAGKTTGGC	420
5	GAAGKTGAAC AGKTGACCAT WACTCTGCCM AATATAGAAA GTTGAAGGAA GCAGTAAAAT	480
	TCAGTATCGT AAAGAACAAC AGCAACAACA ATGTGGAATT CASCCAGGAC TCCCAATCTT	540
	GTAAAACATT CTCCATCTGA AGATAAGATG TCCCCAGCAT CTCCAATAGA TGATATCGAA	600
10	AGAGAACTGA AGGCAGAAGC TAGTCTAATG GACCAGATGA GTAGTTGTGA TAGTTCATCA	660
	GATTCCAAAA GITCATCATC TTCAAGTAGT GAGGATAGTT CTAGTGACTC AGAAGATGAA	720
15	GATTGCAAAT CCTCTACTTC TGATACAGGG NAATTGTGTC TCAGGACATC CTACCATGAC	780
	ACAGTACAGG ATTCCTGATA TAGATGCCAG TCATAATAGA TTTCGAGACA ACAGTGGCCT	840
	TCTGATGAAT ACTTTAAGAA ATGATTTGCA GCTGAGTGAA TCAGGAAGTG ACAGTGATGA	900
20	CTGAAGAAAT ATTTAGCTAT AAATAAAAAT TTATACAGCA TGTATAATTT ATTTTGTATT	960
	AACAATAAAA ATTCCTAAGA CTGAGGGAAA TATGTCTTAA CTTTTGATGA TAAAAGAAAT	102
25	TARATTIGAT TCAGARARA ARARARARA ARCTCGA	105

(2) INFORMATION FOR SEQ ID NO: 205:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 721 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 205:

GAATTCGGCA CGAGTCATCC CTCTCCCTCT TTCACTCCCT TACTCTTACT CTGTTTTTTG 40 TOCTCCAGAC AGACAGACCC TACCTCTTTT GCTTCTTTT TGTTTGTTTG TTTTGAGATG 120 GAGTGTCGCT CTTGTTGCCC AGGCTGGAGT GCAGTGGCGC AATCTCGGCT CACCACAACC 45 TCTGCCTCCC GGGTTCAAGC AATTCTCCTG CCTCAGCCTC CCGAGAAGCT GGGGATTACA 240 GGCATGCGCC ACCACACCCA GCTNAATTTT ATATTTTTAG TAGAGATGGT GTTTCTCCAT 300 GTTGGTCAGG CTGGCCTCAA ACTCCCAACC TCAGGTGATN CCGCCTGCTT TGGCCTCCCC 360 50 AAAGTGCTGG GATTACAGGC GTGAGCCACT GCGCCCAGCC TCTTTTGCTC CTTTATACTC 420 ATTAACTCAC GCCTGTAATC CCTGTTTTGG GAGGCCAAAG TGAGAAGGTT GCTTGAGGCC 480 55 AAGAGTTTGA GACTAGCCTG GGCAACACAG CAAGATGCCA TCTTTATAAT AAAAATAAAA 540 ATAAAAATCA ATTAGCTGGG CATGGTGGAA CGCACCTGTA GTCCCAGCCA ATTGAGAGGC 600 TGAAGTGGGA GGATCATTGA GCCCAGGAGT TGAGGTTGCA GTGAGCCATG ATCATGTCAC 660 60

458

	TACACTCAGC CTGGGCAATA GAGGGACATG TTGTCTCTAA AAAAAAAAAA	720
	A	721
5		
0	(2) INFORMATION FOR SEQ ID NO: 206:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2465 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
15	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 206:	
	CCACCATTTA TCCAACTGAA GAGGAGTTAC AGGCAGTTCA GAAAATTGTT TCTATTACTG	60
20	AACGTGCTTT AAAACTCGTT TCAGACAGTT TGTCTGAACA TGAGAAGAAC AAGAACAAAG	120
	ACCCAGATGA TAAGAAAGAG GGAGGTAAAG ACAGAGCTTT GAAAGGAGTT TTGCGAGTGG	180
25	GAGTATTGGC AAAAGGATTA CTTCTCCGAG GAGATAGAAA TGTCAACCTT GTTTTGCTGT	240
	GCTCAGAGAA ACCTTCAAAG ACATTATTAA GCCGTATTGC AGAAAACCTA CCCAAACAGC	300
	TIGCIGITAT AAGCCCIGAG AAGIATGACA TAAAATGIGC TGTATCIGAA GCGGCAATAA	360
30	TTTTGAATTC ATGTGTGGAA CCCAAAATGC AAGTCACTAT CACACTGACA TCTCCAATTA	420
	TTCGAGAGA GAACATGAGG GAAGGAGATG TAACCTCGGG TATGGTGAAA GACCCACCGG	480
35	ACGTOTTGGA CAGGCAAAAA TGCOTTGACG CTCTGGCTGC TCTACGCCAC GCTAAGTGGT	540
	TCCAGGCTAG AGCTAATGGT CTGCAGTCCT GTGTGATTAT CATACGCATT CTTCGAGACC	600
	TCTGTCAGCG AGTTCCAACT TGGTCTGATT TTCCAAGCTG GGCTATGGAG TTACTAGTAG	660
40	AGAAAGCAAT CAGCAGTGCT TCTAGCCCTC AGAGCCCTGG GGATGCACTG AGAAGAGTTT	720
•	TIGARISCAT TICTICAGGG ATTATICTTA AAGGTAGTCC TGGACTICTG GATCCTTGTG	780
45	AAAAGGATCC CTTTGATACC TTGGCAACAA TGACTGACCA GCAGCGTGAA GACATCACAT	840
	CCAGTGCACA GTTTGCATTG AGACTCCTTG CATTCCGCCA GATACACAAA GTTCTAGGCA	900
	TOGATCCATT ACCOCAAATG AGCCAACGTT TTAACATCCA CAACAACAGG AAACGAAGAA	960
50	GAGATAGTGA TGGAGTTGAT GGATTTGAAG CTGAGGGGAA AAAAGACAAA AAAGATTATG	1020

ATAACTITTA AAAAGTGTCT GTAAATCTTC AGTGTTAAAA AAACAGATGC CCATTTGTTG

CATGGAAGAA CCAAGTTTTT CTATGATATT AAAAAATGTA CAGTGTTAGG TATTATTTGA

ATGGAAAGAC ACCCAAAAAA AAAAATGTGC TCCGACTAGG GGGAAAACAG TAGTTCCGAT

55 GCTGTTTTC ATTCATAATA ATGTCTACAT TGAAAAATTT ATCAAGAATT TAAAGGATTT

1080

1140

1200

1260

	TTTTTCCCAT TATTTTATT TTATTTTCTG GTTGCCCTAG CTTCCCCCCC TATTTTTGTG	1320
	TCTTTTATTA ACTAGTGCAT TGTCTTATTA AATCTTCACT GTATTTAATG CAGGATGTGT	1380
5	GCTTCAGTTG CTCTGTGTAT TTTGATATTT TAATTTAGAG GTTTTGTTTG	1440
	CTAGTTGTAA GTTACTTTGT TATAGATGGT ATCCTTTACC CCTTCTTAAT ATTTTACAGC	1500
10	AGTACGTTTT TITGTAACGT GAGACTGCAG AGTTTGTTTT TCTATATGTG AAGGATTACA	1560
10	ACACAAAAAG TTATCCTGCC ATTCGAGTGC TCAGAACTGA ATGTTTCTGC AGATCTTGTG	1620
	GCATTTGTCT CTAGTGTGAT ATATAAAGGT GTAATTAAGA CAGAGTTCTG TTAATCTAAT	1680
15	CAAGTITICCT GTTAGTIGTG CATTAGCAGT ATAAAAGCTA ATATATACTA TATGGTCTTG	1740
	CAACAGTITT AAAGCCTCTG CATAATIGAT AATAAAAATG CATGACATTC TIGITTTTAA	1800
20	TAGACTITTA AAATCATAAT TITAGGTTTA ACACGTAGAT CTTIGTACAG TIGACTITIT	1860
20	GACATAGCAA GGCCAAAAAT AACTITCTGA ATATTTTTTT CTTGTGTATA AGTGGAAAGG	1920
	GCATTITICA CATATAAGIG GGCTAACCAA TATTITCAAA AGAACITCAT CATIGTACAA	1980
25	CTAACAACAG TAACTAGCCC TTAATTATGG TGACAGTTCC TTATTGGTGT GTGTGAGATT	2040
	ACTCTAGCAA CTATTACAGT ATAACACAGA TGATCTTCTC CACACACCCC ATCACCCAGA	2100
30	TAATTTACAG TICTGITAAC AGIGAGGITG ATAAAGTATI ACIGATAAAA AATTATCTAA	2160
30	GGAAAAAAAC AGAAAATTAT TTGGTGTGGC CATCTTACCT GCTTATGTCT CCTACACAAA	2220
	GCTAAATATT CTAGCAGTGA TGTAATGAAA AATTACATCT TACTGTTGAT ATATGTATGC	2280
35	TCTGGTACAC AGATGTCATT TTGTTGTCAC AGCACTACAG TGAAATACAC AAAAAATGAA	2340
	ATTCATATAA TGACTTAAAT GTATTATATG TTAGAATTGA CAACATAAAC TACTTTTGCT	2400
40	TTGAAATGAT GTATGCTTCA GTAAAATCAT ATTCAAATTT AAAAAAAAAA	2460
70	CTCGA	2465

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(2) INFORMATION FOR SEQ ID NO: 207:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1480 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 207:

55
GAATTCGGCA CGAGCTCAAG CTGGCAGGTG GTCGGGGGAG CGGCCGGAGA GGAGCTGCCG 60
GGAGTTCGTG CCCTGCAGGA CATGACACCA GTGGCATATC ACGGCCATGG GGTCTCAGCA 120
60 TTCCGCTGCT GCTCGCCCCT CCTCCTGCAG GCGAAAGCAA GAAGATGACA GGGACGGTTT 180

•		
	GCTGGCTGAA CGAGAGCAGG AAGAAGCCAT TGCTCAGTTC CCATATGTGG AATTCACCGG	240
5	GAGAGATAGC ATCACCTGTC TCACGTGCCA GGGGACAGGC TACATTCCAA CAGAGCAAGT	300
	AAATGAGTTG GTGGCTTTGA TCCCACACAG TGATCAGAGA TTGCGCCCTC AGCGAACTAA	360
	GCAATATGTC CTCCTGTCCA TCCTGCTTTG TCTCCTGGCA TCTGGTTTGG TGGTTTTCTT	420
10	CCTGTTTCCG CATTCAGTCC TTGTGGATGA TGACGGCATC AAAGTGGTGA AAGTCACATT	480
	TAATAAGCAA GACTCCCTTG TAATTCTCAC CATCATGGCC ACCCTGAAAA TCAGGAACTC	540
	CAACTICTAC ACGGTGGCAG TGACCAGCCT GTCCAGCCAG ATTCAGTACA TGAACACAGT	600
15	GGTGAATTIT ACCGGGAAGG CCGAGATGGG AGGACCGTTT TCCTATGTGT ACTTCTTCTG	660
	CACGGTACCT GAGATCCTGG TGCACAACAT AGTGATCTTC ATGCGAACTT CAGTGAAGAT	720
20	TTCATACATT GGCCTCATGA CCCAGAGCTC CTTGGAGACA CATCACTATG TGGATTGTGG	780
	AGGAAATTCC ACAGCTATTT AACAACTGCT ATTGGTTCTT CCACACAGCG CCTGTAGAAG	840
	AGAGCACAGC ATATGTTCCC AAGGCCTGAG TTCTGGACCT ACCCCCACGT GGTGTAAGCA	900
25	GAGGAGGAAT TOGTICACTT AACTCCCAGC AAACATCCTC CTGCCACTTA GGAGGAAACA	960
	CCTCCCTATG GTACCATTTA TGTTTCTCAG AACCAGCAGA ATCAGTGCCT AGCCTGTGCC	1020
30	CAGCAAATAG TIGGCACTCA ATAAAGATTT GCAGAATTTA ATACAGATCT TITCAGCIGT	1080
	TCTTAGGGCA TTATAAATGG AAATCATAAC GTGGTTCTAG GTTATCAAAC CATGGAGTGA	1140
	TGTGGAGCTA GGATTGTGAG TGACCTGCAG GCCATTATCA GTGCCTCATC TGTGCAGAAG	1200
35	TCGCAGCAGA GAGGGACCAT CCAAATACCT AAGAGAAAAC AGACCTAGTC AGGATATGAA	1260
	TYTGTTTCAG CTGTTCCCAA AGGCCTGGGA GCTTTTTGAA AAGAAAGAAA AAAGTGTGTT	1320
40	GCCTTTTTT TTTTTTAGAA AGTTAGAATT GTTTTTACCA AGAGTCTATG TGGGGCTTGA	1380
٠.	TYCACCCTTC ATCCATTGGC TGGAACATGG ATTGGGGATT TGATAGAAAA ATAAACCCTG	144
	CTTTTGATTC AAAAAAAAA AAAAAWAAA AAAAACTCGA	148
45		

(2) INFORMATION FOR SEQ ID NO: 208:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 872 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 208:

CAGTATTICC CTCAGTACTG TAAGCAAAAG TGGTATGTTT TTCTTTCTTT ATGTCTACTC

461

	TGTCCTCTGT GGCCTTCTGG TGTACCCCTC TCTTCCTAGC CATTCAGTCT CTCTAGTCAC	120
	CTCCCTAGTA GCTAGTGCTC TCTAAGTTTT TATTTAATTA GAACAACTCC ATTTCCATTT	180
5	CAAGGTAGGT CAATGGGGG AAAAGCCTCA TGATTTAAAC TGAAGTTAAC AACACAGCTT	240
	TTAAAATGAA AACTCATACT CCAACTTCTA AAGTATATTT GAGCTGATTT GTTTCCAAAA	300
	CAAAGATATG CTGTACCTAA AACTGCTAAA ACAAAAATAT AAAGACAAGG ACTAGGTGAT	360
10	TAAGGGGAGA GAAAAATCAT YTCTTTTCCA GGAAACCTTT GCTAAAATAA GCAAAACTTG	420
	ANTICTATGCT TCATGGAAAC TGACACAAAG AAAAGAAACT GATGGATTGC ACAGGCCTTG	480
15	TTATAGAAAT AGATCTATAA AAAGATCTGT CCACAGGAAA TATACACCTT CTCCTGGTTC	540
	TGAACTICAA TGGGGATTIG TCACCTAGGT CICCATCTAT AGGAATACCT TCACATACCT	600
•	ATCTATTCAT GCACATATTC TGAAAACAGG TACATACAAA ATTACAACAA AGGAAAAAAA	660
20	TTCTATTGAA CACTTAAAAA TAGAAACAGG CCAGGCACGG TGGCTCATGC TGTAATCCCA	720
	ACAATTTGGG AGGCTGAGGC TGGTGGATCA CCTGAGGTCA GGAGTGTGAG ACCAGCTTGG	780
25	CCAACATGGT GAAACCCCGT CACTACTAAA AATACAAAAA AAATTAGCCT GTGTGGTGGC	840
	ACACTCNTAC AATCCNGGCT GACTCGGGAA AN	872
30		
50	(2) INFORMATION FOR SEQ ID NO: 209:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1779 base pairs (B) TYPE: nucleic acid (C) STRANDEENESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 209:	
	AATTGCCAAG ACTGCACAAA ATTACAGTGC TAATGTATAT GGTTGCAGTT CACATAAAGA	60
45	CAAAAGCATC TGTTATGAAA TGAGTAGTAA TATTGGGTGG TTGATTTGTT CTTAGCAGAC	120
43	TIGGCITCAT WITGGICITG AGATAAAATG GCCAGCATAA AIGCIGITTA TATICACGIT	180
	TTCCTAGGTG TGTGTGCA GGCCACAGCA GCATGCCCTT GGTGTAGTCA GTGCCGAAAS	240
50	GGGTCTGTTC CTTCTTGAGC CTGCCTGCAG GGATGGTCTC CTTTTAAAGC AGGTTGTGTG	300
	CAGCATTCAG TACACTGAAG GTAAGCTAAA CCATCAACAT CTCTGGTGTT TTAAGATGTT	360
£ £	ATTITATIOG AACAACIGAC AAAIGAGGGA IGITAGCITI GIGGCAGAAI ICCCIGCAIG	420
55	TGTGATAACT GATCTTGTTT TATTTTTTGG CATTGCAACT GTGGCATAGT TACAATTTCT	480

GTTTGKTCAT CACATTTAAA ATTGGRAGAG AACGCGCTTG AKGGATAGAG CGCCTTCAGK

60 GTACTGTTC TTATTAACIT TACTTTTTT AAATCAACTT GCTATAGACT TTATATACAT

540

-	TITGITAAAT ATAGITCCTA GIGACATAGA AACGATGCGT AGITTICATT TACTAATTAC	660
		500
5	AAATGTTGAG GCCTAATTCT GAAAGTCCTC ATATTTAAAG GCTAGACAAC GTAATGAAAT	720
_	TTTTAACTAT TTGTATGTCA TTTTGAAAGT GTACTGCTTT ATGGTAAAAG TGTTTTTCAT	780
	TIGITCATTG TITICATTAT TIGIGATCAT GITGICTITC AATACAGGCA TAAACCTICC	840
10	ACTOTTGAAC AAAGCAGCTG CTTTTTAAAA GCGGTAATTG CTTCTTTACC TTTTATTTCT	900
	TYTGTAAATG AAGCTYTTCT TTAAGAATGT GACTYTAAAG TGTTGTCTAT TGCATAAAAC	960
15	AGITGACACT CACTTATTGT AAAGTGAAGA TTGTTCTACT GCATGTGAAG TGGACCATGC	1020
13	AGATTTCTGT ATGTTCTCAG TATGCATCAC TAGATAATAA AGTCTTTTGT GAACAAGGCA	1080
	TTTGTAGCCA TTTTTAAAAG TTTTTGTCTT CAGTGCTGGT AAGTCAGGTA AACCATAAAT	1140
20	AGTTAAAAGC AACCTTTIGT TTTTTTCCTG AAAGTTTTTA ATTGAAAGTA TTATTAGTTA	1200
	AAGATGTAAA CCTAGCCAAA ATTACCAGTT TATTAATAAT TAGGATCCTA ATTATTTCAA	1260
25	AAAATCCTAC AAATATTGTC AGCTTTCAGT GTAGTGAGAT TATTCCTGTA GGTTATGGGG	1320
23	TATAATTCAG GATTTAACTA ATGTTTCTGC TATTTTCTCA CTTTTCCTTT TGATGGTGCG	1380
	GAAAGAGAAA AAGGAAAACG GGGCACAGGC CATTCGACGC CTTCTCCAAG GGGTCTGATT	1440
30	TGCTGAGACA CCAGCTTCAC CTTCTTAACA AGGCACCTAA TTACAACAAG CATGCACATT	1500
	TTGGTGCATT CAAGAATGGA AAATCAGAAT AGCAGCATTG ATTCTTCTGG TGCAGCTCAG	156
35	TGGAAGATGA TGACAACCAG AAGACATGAG CTAAGGGTAA GGGACTGTTC TGAAGAACCT	162
))	TTCCATTTAG TGATCAAGAT ATGGAAGCTG ATTTCTGAAA ATGCTCAGTG TGTACTCTAA	168
	TTATTTATGG TACCATTIGA ATTGTAACIT GCATTTTAGC AGTGCATGTT TCTAATTGAC	174
40	TTACTGGGAA ACTGAATAAA ATATGCCTCT TATTATCAA	177

45 (2) INFORMATION FOR SEQ ID NO: 210:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2110 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 210:

	GCCTCGGAGG GCCCTCGGCT GCCCCACCCT CGGAGCCACT GCTAGAAGGG GCCGCTCCCC	240
	AGCCTTTCAC CACCTCTGAT GACACCCCCT GCCAGGAGCA GCCCAAGGAA GTCCTTAAGG	300
5	CTCCCAGCAC CTCGGGCCTT CAGCAGGTGG CCTTTMAGCC TGGGCAGAAG GTTTATGTGT	360
	GGTACGGGGG TCAAGAGTGC ACAGGACTGG TGGWGCAGCA CAGCTGGATG GAGGGTCAGG	420
10	TGACCGTCTG GCTGCTGGAG CAGAAGCTGC AGGTCTGCTG CAGGGTGGAG GAGGTGTGGC .	480
	TGGCAGAGCT GCAGGGCCCC TGTCCCCAGG CACCACCCCT GGAGCCCGGA GCCCAGGCCC	540
	TOGCCTACAG GCCCGTCTCC AGGAACATCG ATGTCCCAAA GAGGAAGTCG GACGCATGGA	600
15	AATGGATGAG ATGATGCCGG CCATGGTGCT GACGTCCCTG TCCTGCAGCC CTGTTGTACA	660
	GAGTCCTCCC GGGACCGAGG CCAACTTCTC TGCTTCCCGT GCGGCCTGCG ACCCATGGAA	720
20	GGAGAGTGGT GACATCTCGG ACAGCGGCAN CAGCACTACC AGCGGTCACT GGAGTGGGAG	780
20	CAGTGGTGTC TCCACCCCCT CGCCCCCCCA CCCCCAGGCC AGCCCCAAGT ATTTGGGGGA	840
	TECTITIEGT TCTCCCCAAA CTGATCATEG CTTTGAGACC GATCCTGACC CTTTCCTGCT	900
25	GGACGAACCA GCTCCACGAA AAAGAAAGAA CTCTGTGAAG GTGATGTACA AGTGCCTGTG	960
	GCCAAACTGT GGCAAAGTTC TGCGCTCCAT TGTGGGCATC AAACGACACG TCAAAGCCCT	1020
30	CCATCTGGGG GACACAGTGG ACTCTGATCA GTTCAAGCGG GAGGAGGATT TCTACTACAC	1080
50	AGAGGTGCAG CTGAAGGAGG AATCTGCTGC TGCTGCTGCT GCTGCTGCCG CAGACCCCCA	1140
	GTCCCTGGGA CTCCCACCTC CGAGCCAGCT CCCACCCCCA GCATGACTGG CCTGCCTCTG	1200
35	TOTGCTCTTC CACCACCTCT GCACAAAGCC CAGTCCTCCG GCCCAGAACA TCCTGGCCCG	1260
	GAGTCCTCCC TGCCCTCAGG GGCTCTCAGC AAGTCAGCTC CTGGGTCCTT CTGGCACATT	1320
40	CAGGCAGATC ATGCATACCA GGCTCTGCCA TCCTTCCAGA TCCCAGTCTC ACCACACATC	1380
40	TACACCAGTG TCAGCTGGGC TGCTGCCCCC TCCGCCGCCT GCTCTCTMTC TCCGGTCCGG	1440
	AGCCGGTCGC TAAGCTTCAG CGAAGCCCCA GCAGCCAGCA CCTGCGATGA AATCTCATCT	1500
45	GATCGTCACT TCTCCACCCC GGGCCCAGAG TGGTGCCAGG AAAGCCCGAG GGGAGGCTAA	1560
	GAAGTGCCGC AAGTGTATGG CATCGAGCAC CGGGACCAGT GGTGCACGGC CTGCCGGTGG	1620
50	AAGAAGGCCT GCCAGCGCTT TCTGGACTGA GCTGTGCTGC AGGTTCTACT CTGTTCCTGG	1680
30	CCCTGCCGGC AGCCACTGAC AAGAGGCCAG TGTGTCACCA GCCCTCAGCA GAAACCGAAA	1740
	GAGAAAGAAC GGAAACACGG AGTTTGGGCT CTGTTGGCTA AGGTGTAACA CTTAAAGCAA	1800
55	TTTTCTCCCA TTGTGCGAAC ATTTTATTT TTAAAAAAA GAAACAAAAA TATTTTTCCC	1860
	CCTAAAATAG GAGAGAGCCA AAACTGACCA AGGCTATTCA GCAGTGAACC AGTGACCAAA	1920
60	GAATTAATTA CCCTCCGTTT CCCACATCCC CACTCTCTAG GGGATTAGCT TGTGCGTGTC	1980
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	AAAAGAAGGA	ACAGCTCGTT	CTGCTTCCTG	CTGAGTCGGT	GAATTCTTTG	CTTTCTAAAC	2040
	TCTTCCAGAA	AGGACTGTGA	GCAAGATGAA	TTTACTTTTC	TTAAAAAAAA	АААААААА	2100
5	AAAAACTCGA	•					2110

10 (2) INFORMATION FOR SEQ ID NO: 211:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 938 base pairs

(B) TYPE: nucleic acid

15 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 211:

20	GGCACAGGAA AAAAAAGAAA AAAGAAAAAA GAAAAAAGTT TTTGTACCCA CAGATTAGCA	60
	TTTTCTTGAT GTTTGAAAAA AGTTTAAGCT ATGTCCTAAT TTAAAAATGA GCACAAACTA	120
	CTTAACAGAT GTCTGTTCCC TCTTCTCTTA CTTAAATTAT CTTTATTTTC ACCATCACCT	180
25	CCCAGTGCCG AACACCTGAN CTCTGTGTTT TGTGGTTGGA TCCTGGGTTG CCAAGTTCCT	240
	ATTTOGTCAG TCCCTGGCCT GTGGGGCGGT CTCAGGAAGT GGCATGCTCT TCAMGRAGGA	300
30	TOGITCATYI CCAGIATAAC CAWITIGITA ATAATAGIIG ATAATICCCA GCITITACCA	360
	GATGARTITT GACTTATTIT TCCTCCTTTG ACCTGTTCAA AGCTAACATA TCTCGGTCAG	420
	TTCCGAGAGG GTGGGGGATT TGAGAATGTG AGGAGGAGTG GGGTTAGAAT GGGTTTGCCT	480
35	ATCTGGGCAA GGAAAGAGTT CCTAGTCGAT TGGGCACAAT GACAAAATGA TTCCATGGAT	540
	AGAATCGTCC CATGTTGCTG GAACACCTCA CGTGTTGTGA ACGCCTTAAA TTCCTGCCAT	600
40	CCCTTCTCTG ATTCCCCACC TCCCTGTAGT TTCCACAGGA TTTATCTCTC TGTACCCCCG	660
40		
45	TCCTCCAACT CTACTCTGTC AGCCTCTCCT CCATCCCTTA CTTCCCTTCT AAATTCCAGG	720
	AGATGACCIC ACTITGCAAA GCAAATTGGA GCCACCAAAT TGTAGCTCTC CTCGGTGGAA	780
	ACTGCATCTG TGCTCATCCC TGCACCTTCT TGCAGAAAGC CGCCCCTCA GGCCAAGATG	840
	AGTGCCTGGC CCCCATGGGA GACTCAGACA CTTTGACCCC TTGTGACTTC AGCATCTCCC	900
50	TCTTTAAAGA TTCTCTCCCA ACATTCAGTC GTGCTCGA	938

55 (2) INFORMATION FOR SEQ ID NO: 212:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1551 base pairs

(B) TYPE: nucleic acid

60 (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 212:

5	AGCTGGACT AAGCATAGAG AACCAGGAGA GAAAGAAAGA TTTAAGAGAC TGAGTAATAT	50
•	TTTTTGACAG ATCATTTAAG AAACTGAGTA ATTTTTTTT TCTCCAAAAG GGCATSGGTT	120
10	THITTITIGT THIGHTHIT CICIATURG CACTURETAG GGATIGGICT ATAAATTITT	130
	TGAAAGATCA TAGGATAAAT TTCTTTGTAG CAACTTCCTA TTTTAGTGTT TATGTTAGGG	240
	GARCCCCARG TOTCCCTGCT GATACGCCAT TAGGGCCACT TCTCAGCCTC TGGCTACATC	300
15	ATAATGCTTT TTTTTCTATC TTGCCAAAGT TTCCMGAAAA TTKAKGTTTT CTAATTTAA	360
	AAAAATTGGT TGTGGAGATG GGATGGGACC TCTTTATAAG CCCTGAAAAT AAGTGATTTN	420
	TTTTAAGTGC TATTCTGCTA TAAACCTGAT TCTCACTTTT TTCTGTAGAC AACAGTTTTT	430
20	TATAATATAT CTATTTTGTG TGGACATTAT TTCCTTTTAA CCAATACTGA AATTCCATAG	540
	TGTAWACTTT CTCCACATTT TCTTTGATTA ATACTTYCTT AAAATAGACA CTTGGATTGG	630
25	CACCAGCTGT CACCAATAAA GCTGCCCTGA ACATTGTCAA TCAATCCTGT TAACCAATTT	650
	GAGAATTITT CTGGAATGCT TAGTTAGGGA TGAAATTGCT GGGTTATAGG TATGAGTATG	720
	CTTGATATAC TTTTCTCCAG AATGTCTACA CCTGTGTGTA CACCACATCT CCAGAGATAG	730
30	GGGAATCTTA TGTCCCTGCT AACTGCTCTC GTTATTTAAT TTTCTGACAT TTGCCGCCGC	840
	COCCCCCCC TOCCCCCAAC ACACACATGG TATAAAGTGG TAGTTTCTTG TTTTAAATTG	900
35	AACTITIGAA TGATTIGAAT TIGGGCATIT CTITGTATCC TGAGTTATTT TGGTTTCCCG	950
	TTATGTGAAT ATCCTTTICC TATGCTTTAA CTACTTTTCT AATTTGTCCC TTTTTTNGGT	1020
	TATCABATTC CAGGCCATTG TCTATTCCAT CGTCACTTTT GGGTATTGGA BACATCTTTC	1030
40	CATTCTGTAG CCTGTCTGTT GAACATAAAT CTTGATTTTT ATGTAATCAG ATTTTTCTCC	1140
٠.	TTACCGTTAT GTTCTTGGAA TTTTATTTAA GAAATCTTTT TCTATCCTGA GACCACAAAA	1200
45	TATCHTAAGGC	1260
	ATGTGTAGTT CATTITATAT GGTGTGAAAT AGTTCTTATT CATTTATTCA ACACATATTG	1320
50	GTGGAGTGCC TGCTGATGGT AGTACTCTTC AGAGTACTTT GTATATATTT GTGAACACAT	1330
	ATTCTTGCCC TGGAAGCTTA TGTTGTCNTT CAAGGTAGAT CCNTACTCGG TTTCCACCTG	1440
	TITTCTTCAG CCCTCAGGAT GAATTCCACA ATTTTACACA TAGCACCAGT TAAGGAATAG	1500
5:		1551
J.	· ++++	

5	(A) LENGTH: 997 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 213:	
10	AGAGAGTOCT CAACAGAACC TAATCATGCT GGCACCOTAA TOTCATACTT CTAGCOTCCA	50
	GAACTGAGAG AACATAAACT CCAGTTGTTT AAGCTACCCA GECTATGGTA TTTGTTATTA	120
15	TAGCCCAAGC TAAGTCAGGT GGAAAGGCAG AAATATTTTG AGAAGARTCA TTTCTACAAA	130
	AACAGAGTIG TICTAAATGA AATGGCCAGA TATTICATC: TOTTCATACT AGTATTIATG	240
	ARAGITICAT TARACACCAC TIGGCCAGCA CCCAGGCCIG CCACCCTICAG AACGGCAAAC	300
20	AAAAGCAAAT GATTTGAGGA ACAAAAGAGT GGACACAGAG CTTCTCAGAA GATGGCTCCA	360
	TOTTOTGAGA TGATOTTOTG AGATCATCAA TTTTOTGCAC CTGATGTOOT ACTOCAALTG	420
05	TAGTAGATAA GAGCAAAGAC ACTTCCTGAT CCTGTGGAAA ATGCTGGAGC CCTGCTGATG	430
25	GAGAGGCTGA CACTGGGACC AACAGAAGGC CGGACATTTA TYTGCTGCAG CCCTTCTGCA	540
	CCTGGGCCCT CTTCAGGCCT TGTACCTTGC ACTCCCCATG CCACTGCAGC ACCTGGTAAG	600
30	CTGAAGTTAG GTATTTGAAG AGATAATTTG CCCCCAACAA AGAATTACTT AAAAGAAAAA	550
	GGAAACCACT AAATTCCACT TGACAAACCA GTTTGTTCAG TTTTTACTTT TGCAAATTTG	720
35	ARACTITCTC TITGGCACCA TATGATTCTG TTACATTAGG GCTCATCAAT GCTAAGATAC	780
33	ACAGCTAGGT CTACCAGCTG CCAGTGGTCA AGAATGAAAG AACCTCTCAG AGAGAGATCA	240
	GTTTCTAATA ACCTAACAGT TTTCCTTGGS TATTACMAAA ARAAAAAAA TTAGAATAAA	900
40	ATGTCAGTGC CATGCAGGCA AGTACAGATA TGGAAATGAA AGCTTTGTCT ACAACTGCAA	960
٠,	GATTIGITIG TTAATAAAAT TGATTGGGAT CACTCGA	997
45		
	(2) INFORMATION FOR SEQ ID NO: 214:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1496 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 214:	
	GAATTOGGCA CGAGTGACCA CAGATATOTT TGGCTTTCAG CCTCACCACA ATGCTGTCCA	60
60	CTATGTTTT TTTAATCGAT TGACATCTCA TGAATCCACA AATTTAGCCG CTTTTCCATC	120

	TITTCCATCT TIGTCATAGC TICATCACGC ACGATGGAGG TCACTTCAGC ACTATCCGGA	180
	GCGGCCTCAC GGACAGATCR GTGAATTTCC TTTTCCTTTT TCTTGATGTA CCGGATTGTC	240
5	GACTOGITAA CATIGAGOTO ATGGOCCAACA GCACTGTAAC TCATGCCTGA TTGGAGOTTA	300
,	TOCAACACGC GGAMTTTCTC CGTAAGGSAM ATCAMGGTCT TCTTTCGCTT AGGAACACTG	360
10	GGCARARCTT AARCACTACG CTTGGGGGCC ATTTTAGAAA GCAAAACCAC CCACAAAAAG	420
	CAGAAAAAA AGTGTCAGTA AACAGACTGN NGANAGGACT CTTTGTTTAC AGCACAGGAG	480
	CTGCGACTAG AAGGCGGCGC TTCTCCCCAG TTCAAACTTC AGCTGGGAAC CTTACCTCCG	540
15	CCAACTCCAA ATTITCACCC TCTGCGCATG CCCGGGAAAS AAACCCCCAG AACAGTACCG	600
	TGATGATTGA TTTTAGGGTT ACAAATACAT TTTAGCAAGT AAGTGAATTT GGCATTACGA	660
	ATTAATGATT AATGAAGGTC ACCIGIATIT CCATAGATAT GTAATTITAT TTAAGCAGGT	720
20	TTATTATATT AAGGCGGGGA GGCAGCGCCG AAGACTACAA GTTCCAGCAT GCACCGCGTC	780
	CGGGCGGGTT CGGGCTCCCA GCGAGGGCTT CAGGGACGCC AGCCCGGAGG CATCGGCCGG	840
25	AAGTGTCGTA GGGCAACCAC GTAGTACTCT CTGCGCATGT GCAAAGCGCT GTCGGGGGCC	900
	GCCCTAGCTG CCGTCGCCGC CGCCGGGCT CTATGGTCTC TCCCTAGAGC TTTGCCGTTG	960
	GAGGCGGCTG CTGCGGTCTT GTGAGTTTGA CCAGCGTCGA GCGGCAGCAA CATGGAGGAA	1020
30	TTCGACTCCG AAGACTTCTC TACGTCGGAG GAGGACGAGG ACTACGTGCC GTCGGGTGAG	1080
	CGATTCCGCC TGAGGCGAGA AGCGAATTGC CCCGCCCCAC GCCTCACGTG AGGCGCGCTC	1140
35	TECCCCCCCC GGCGTCTGCC CTGTGGCCCCA GGTGGTCCAG GGGGGCTCCT GTTCTCGAGC	1200
	GTCCGCTCCC TCAGGCCCCT CATQCTCGGC CGCTCCGGCC CGAGGCGTGT GCGCGTGGCG	1260
40	GITCTGTGCT CCCCTCCCGT TGGGCAGCTC CGGCCGCCGC CCCCTCTTGC AGCGCGGGAA	1320
	CGGCACATGG ACACGGCCCC TTGTCGCTAG GGACGCTCGT CGGTCAGCCC CGAACGACAA	1380
٠.	CGCTGCTTCA GAAGTCGGGG CGGCAGTTCG AGCCTTGGAA GTTTTTTTCA GCCCTGGCCC	1440
45	CAMPAGNACING TOTAL TOTAL CONCINCTION CONCINCING	1496

50 (2) INFORMATION FOR SEQ ID NO: 215:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1308 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 215:

	CIGCCITTGA CCCATCACAC CCCATTTCCT CCTCTTTCCC TCTCCCGCT GCCAAAAAAA	120
5	AAAAAAAAGG AAACGTTTAT CATGAATCAA CAGGGTTTCA GTCCTTATCA AAGAGAGATG	180
	TGGAAAGAGC TAAAGAAACC ACCCTTTGTT CCCAACTCCA CTTTACCCAT ATTTTATGCA	240
	ACACAAACAC TGTCCTTTTG GGTCCCTTTC TTACAGATGG ACCTCTTGAG AAGAATTATC	300
10	GTATTCCACG TTTTTAGCCC TCAGGTTACC AAGATAAATA TATGTATATA TAACCTTTAT	360
	TATTGCTATA TCTTTGTGGA TAATACATTC AGGTGGTGCT GGGTGATTTA TTATAATCTG	420
1.5	AACCTAGGTA TATCCTTTGG TCTTCCACAG TCATGTTGAG GTGGGCTCCC TGGTATGGTA	480
15	AAAAGCCAGG TATAATGTAA CITCACCCCA GCCTTIGTAC TAAGCTCTTG ATAGTGGATA	540
	TACTOTTTTA AGTTTAGCCC CAATATAGGG TAATGGAAAT TTCCTGCCCT CTGGGTTCCC	600
20	CATTTTACT ATTAAGAAGA CCAGTGATAA TTTAATAATG CCACCAACTC TGGCTTAGTT	660
	AAGTGAGAGT GTGAACTGTG TGGCAAGAGA GCCTCACACC TCACTAGGTG CAGAGAGCCC	720
25	AGGCCTTATG TTAAAATCAT GCACTTGAAA AGCAAACCTT AATCTGCAAA GACAGCAGCA	780
25	AGCATTATAC GGTCATCTTG AATGATCCCT TTGAAATTTT TTTTTTGTTT GTTTGTTTAA	840
	ATCAAGCCTG AGGCTGGTGA ACAGTAGCTA CACACCCATA TTGTGTGTTC TGTGAATGCT	900
30	AGCTCTCTTG AATTTGGATA TTGGTTATTT TTTATAGAGT GTAAACCAAG TTTTATATTC	960
	TGCAATGCGA ACAGGTACCT ATCTGTTTCT AAATAAAACT GTTTACATTC ATTATGGGGT	1020
35	ATGTATGACC TTCATTTTCC AAGAAATAGA ACTCTAGCTT AGAATTATGG ATGCTCTAAA	1080
33	ATGTCAGAAT GGGAACTCTC CTCGAAGTTC TCCCAAACTC AGAGACAGCA CTGCCTTCTC	1140
	CTARATGATT ATTCTTTTCT CCCTGTTTTC TGGTATTTTC TAGGCATCCT TCTCACCACA	1200
40	GCCATAACCC TTTTTTACTT CCATTAGGCC GTATAACTGG NGGGACNGCT GGTCGGTATA	1260
٠.	TAATACTGGT WCCAACAMAG GGGTTCTGGA TGTACACMAG GTTATCTT	1308
45		
73	216.	
	(2) INFORMATION FOR SEQ ID NO: 216:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1705 base pairs	

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 216:

TOGCCATGGA AGCGCTAGAA GGTTTAGATT TTGAAACAGC AAAGAAGGAT TTCCTTGGAT 60
CTGGAGACCC CAAAGAAACA AAGATGCTAA TCACCAAACA GGCTGACTGG GCCAGAAATA 120

	TCAAGGAGCC CAAAGCCGCC GTGGAGATGT ACATCTCAGC AGGAGAGCAC GTCAAGGCCA	100
	TCGAGATCTG TGGTGACCAT GGCTGGGTTG ACATGTTGAT CGACATCGCC CGCAAACTGG	240
5	ACAAGGCTGA GCGCGAGCCC CTGCTGCTGT GCGCTACCTA CCTCAAGAAG CTGGACAGCC	300
	CTGGCTATGC TGCTGAGACC TACCTGAAGA TGGGTGACCT CAAGTCCCTG GTGCAGCTGC	360
	AGTGGAGACC CAGCGCTGGG ATGAGGCCTT TGCTTTGGGT GAGAAGCATC CTGAGTTTAA .	420
10	GGATGACATC TACATGCCGT ATGCTCAGTG GCTAGCAGAG AACGATCGCT TTGAGGAAGC	480
	CCAGAAAGCG TTCCACAAGG CTGGGCGACA GAGAGAAGCG GTCCAGGTGC TGGAGCAGCT	540
15	CACAAACAAT GCCGTGGCGG AGAGCAGGTT TAATGATGCT GCCTATTATT ACTGGATGCT	600
	GTCCATGCAG TGCCTCGATA TAGCTCAAGA TCCTGCCCAG AAGGACACAA TGCTTGGCAA	660
	GTTCTACCAC TTCCAGCGTT TGGCAGAGCT GTACCATGGT TACCATGCCA TCCATCGCCA	720
20	CACGGAAGAT CCGTTCAGTG TCCATCGTCC TGAAACTCTT TTCAACATCT CCAGGTTCCT	780
	GCTGCACAGC CTGCCCAAGG ACACCCCCTC GGGCATCTCT AAAGTGAAAA TACTCTTCAC	840
25	CTTGGCCAAG CAGAGCAAGG CCCTCGGTGC CTACAGGCTG GCCCGGCACG CCTATGACAA	900
	GCTGCGTGGC CTGTACATCC CTGCCAGATT CCAAAAGTCC ATTGAGCTGG GTACCCTGAC	960
	CATCCGCGCC AAGCCCTTCC ACGACAGTGA GGAGTTGGTG CCCTTGTGCT ACCGCTGCTC	1020
30	CACCAACAAC COGCTGCTCA ACAACCTGGG CAACGTCTGC ATCAACTGCC GCCAGCCCTT	1080
	CATCTTCTCC GCCTCTTCCT ACGACGTGCT ACACCTGGTT GAGTTCTACC TGGAGGAAGG	1140
35	GATCACTGAT GAAGAAGCCA TCTCCCTCAT CGACCTGGAG GTGCTGAGAC CCAAGCGGGA	1200
	TGACAGACAG CTAGAGATTT GCAAACAACA GCTCCCAGAT TCTTGCGGCT AGTGGGAGAC	1260
	CAAGGGACTC CATCGGAGAT NAGGACCCGT TCACAGCTAA GCTRAGCTTT GAGCAAGGTG	1320
40	GCTCARAGTT CGTGCCAGTG GTGGTGAGCC GGCTGGTGCT GCGCTCCATG AGCCGCCGGG	1380
٠.	ATGTCCTCAT CAAGCGATGG CCCCCACCCC TGAGGTGGCA ATACTTCCGC TCACTGCTGC	1440
45	CTGACGCCTC CATTACCATG TGCCCCTCCT GCTTCCAGAT GTTCCATTCT GAGGACTATG	1500
	AGTTGCTGGT GCTTCAGCAT GGCTGCTGCC CCTACTGCCG CAGGTGCAAG GATGACCCTG	1560
	GCCCATGACC AGCATCCTGG GGACGGCCTG CACCCTCTGC CCGCCTTGGG GTCTGCTGGG	1620
50) CTGTGAAGGA GAATAAAGAG TTAAACTGTC AAAAAAAAAA	1680
	ANANANA ARABABABA AANA	1705

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(i) SEQUENCE CHARACTERISTICS:

⁽²⁾ INFORMATION FOR SEQ ID NO: 217:

	(A) LENGTH: 999 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 217:	
	AGCAAATCAC CTTAACGATC TOGAATGAAA CTGTGACCAG TGCCGCCCTG GGTGGTTCTG	60
10	GAGAGACTGC CGTCTTCTTG TTTGGCCATA GGTGCTGGGG CCCCGGCTTC AGTCACTGTC	120
	TCAGACAGKA GTCCCGATAA GCAGATCACC AGTCCTCCAC TGTCCTTCCT GTCGGCCTTG	180
	CIGCATGAGA AGATAGCTGC TTCCTCCCTC TITTCCTACA CTGTAAATTA TTGTTTTACA	240
15	ATTGAGTGYC TTAATAATAG TYTACAAATA CTATGTATTT ATGCAAAACT GTTAAAGTTC	300
	TCATCTGTTA TGATTGGATA CTTGGTCTTG TCAGTAGTGG TCAGCATTGG GTTGTGAGCT	360
20	TGTCCTACTC CATACGTGTT TATCCTGCTA TGCATTTTAC ATTGTGTGTT CACATCTATT	420
	CCAAGGAGCC TTGCTAGAAA CAACACTGGC GGTTCCTGCA GGCCAGGCAG GCATTGGCCC	480
	ATGCTGTGTC CCATAGGAGC CAATGGAAAG AACGTAGCTT GGTCTGCTAG CCAGCCGTGG	540
25	COTOGOGGAG GCCAGGCAGC CTCTGCACCA GAGTCCAGCA CCTGCCCATT CCCCAGTCAC	600
	ACAATCATAC TCTTCTTTCA TAGAGATTTT ATTACCACCT AGACCACCCT AGTTTTCCTC	660
30	TCTGTTAGTG TCCTGAGCTC TTTTGCAACA AAATGTAGGT ACAGACCAAT CCCTGTCCCT	720
	TCCCCAATCA GGAGCTCCAC ACCATGAGTT GTTTGGTTTT CCAGAAGCTG CCAGTGGGTT	780
	CCCGTGAATT GCGTTAAGAT ATCGATGATK TTTTTTATTG TTTTTCTTCT TGTTTTTTTA	840
35	AATAATATA TTAAAGGCAG TATCTTTTGT ACTGTGAATT TGCAGTAGAA GATGCAGAAT	900
	GCACTITUT TITACTICIG TIGGIGIGIA TIGIATATAG TGIGIGIGCT TCTIGIGATG	960
40	TARABARA ADADADAC	999
٠.		
45	(2) INFORMATION FOR SEQ ID NO: 218:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 941 base pairs	
	(B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 218: GGCACGAGTA GCATTICATT TAATCTGCAG GTATATTCTC CCAACAGTT ATTGTCATGT	60
5		120
	GATGTCCTCA GCCAAGATTG TRAGGCAGAG AGGAGCTGTC CCAACCTACT ATACCACCGA	180
6	GCCTGGAGAG ATCATATTT TGGTATTAAA CTGGAGTCTC TCCATCCTTC ACATTGTTGA	190

	TGTCCTCTGT AGCAAACCGG AAAAGTCAGT GACAGAAGAT GCCGCTAGCG GTTTGAGCCA	240
	GAGAATGACA GCTCTGGTTT GGAGAAAAGG GCCGGATGGT GGCTCTAGAA AGCCCATCCT	300
5	TOTGCTCTTC TTTTTTCTCC CCCTTATATT GTGCTTTCAT TCATTCATTC ATTCATCAAA	360
	CATTIGITGA GCACCTATTA TGTGTCAAGC TCTGTGCTAG CCTCTGGAAA ACCTGCCCTC	420
	ATGTAGCTCA CTGTGGAGTA GGAGAAACAA TGACTACACT ATGATAAGCA CGGGTTGTCA	480
10	GGGTCTCACA GAGCAGTGGC CCCTCATCCA GACCGATGAG GTCAAAGAAG GCATCCAGGC	540
	GAGGATGGTG TCAGAGCTAA CTGAAGAATG AGAGGGAGCT GCACCASCAG GGGTTGGAAC	600
15	TGAAGGTGGC AGTGCCTGGA GTCTTGATTC CAGCAGAGGG AGAGCAGTCT GTGAAAAGGC	660
10	ACCAAGGGTG GGAGAGGGCA GAGCACATGG AGGAACTTCA GGTAGTTCTG GATGGCSCTG	720
	GGGCAAAGCT AGAGAGGTAA GAAGAATCTA CAAATGTTCC TCGAGTTACA TGAACTTCCA	780
20	TCCCAATAAA CCCATTGGAA ACGAAAAATT TAAGTCAGAA GTGCATTTAA GGCTGGTCCG	840
	AGTAGAATGA TITTTACAAC GAATTGATCA CAACCAGTTA CAGATGTCTT TGTTCCTTCT	900
25	CCACTCCCAC TGCTTCACCT GACTAGCCTT TAAAAAAAAA A	941
43	Controcke 1001101101	

30 (2) INFORMATION FOR SEQ ID NO: 219:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 575 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 219:

TAAGTGGAAT CCCCCGGGT TGCAGGGAAT TCGGCACGAG GCATTCTGAG AAGCTTAAGA 60 40 CATACTITGA AGACAACCCT AGGGACCTCC AGCTGCTGCG GCATGACCTA CCTTTGCACC CCGCAGTGGT GAAGCCCCAC CTGGGCCATG TTCCTGACTA CCTGGTTCCT CCTGCTCTCC 180 45 GTGGCCTGGT RCGCCCTCAC AAGAAGCGGA AGAAGCTGTC TTCCTCTTGT AGGAAGGCCA AGAGAGCAAA GTCCCAGAAC CCACTGCGCA GCTTCAAGCA CAAAGGAAAG AAATTCAGAC 300 CCACAGCCAA GCCCTCCTGA GGTTGTTGGG CCTCTCTGGA GCTGAGCACA TTGTGGAGCA 50 CAGGCTTACA CCCTTCGTGG ACAGGCGAGG CTCTGGTGCT TACTGCACAG CCTGAACAGA 420 CAGTICIGGG GCCGGCAGTG CTGGGCCCTT TAGCTCCTTG GCACTTCCAA GCTGGCATCT 480 55 540 575 CTCGAGGGG GGCCCGTACC CAATTCGCCC TATAA

121	TNIFORMATION	FOR	SEO	ID	NO:	220:

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5	(i) SEQUENCE CHARACTERISTICS:
3	(A) LENGTH: 3018 base pairs
	(B) TYPE: nucleic acid
	(C) STRANDEDNESS: double
	(D) TOPOLOGY: linear
10	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 220:

GCCAGCCTTA CAGGTTTTAC GTGAAATGAA AGCCATTGGA ATAGAACCCT CGCTTGCAAC 60 ATATCACCAT ATTATTCGCC TGTTTGATCA ACCTGGAGAC CCTTTAAAGA GATCATCCTT 120 CATCATTTAT GATATAATGA ATGAATTAAT GGGAAAGAGA TITTCTCCAA AGGACCCGGA 180 TGATGATAAG TTTTTTCAGT CAGCCATGAG CATATGCTCA TCTCTCAGAG ATCTAGAACT 240 TGCCTACCAA GTACATGGCC TTTTAAAAAC CGGAGACAAC TGGAAATTCA TTGGACCTGA 300 TCAACATCGT AATTTCTATT ATTCCAAGTT CTTCGATTTG ATTTGTCTAA TGGAACAAAT 360 TGATGTTACC TTGAAGTGGT ATGAGGACCT GATACCTTCA GCCTACTTTC CCCACTCCCA 420 AACAATGATA CATCTTCTCC AAGCATTGGA TGTGGCCAAT CGGCTAGAAG TGATTCCTAA 480 540 TCCTGATGCT CATGGCAAGG GACAAGCACC CACCAGAGCT TCAGGTGGCA TTTGCTGACT 600 GTGCTGCTGA TATCAAATCT GCGTATGAAA GCCAACCCAT CAGACAGACT GCTCAGGATT GGCCAGCCAC CTCTCTCAAC TGTATAGCTA TCCTCTTTTT AAGGGCTGGG AGAACTCAGG 720 AAGCCTGGAA AATGTTGGGG CTTTTCAGGA AGCATAATAA GATTCCTAGA AGTGAGTTGC TGAATGAGCT TATGGACAGT GCAAAAGTGT CTAACAGCCC TTCCCAGGCC ATTGAAGTAG 840 TAGAGCTGGC AAGTGCCTTC AGCTTACCTA TTTGTGAGGG CCTCACCCAG AGAGTAATGA 900 GTGATTTTGC AATCAACCAG GAACAAAAGG AAGCCCTAAG TAATCTAACT GCATTGACCA 960 GTGACAGTGA TACTGACAGC AGCAGTGACA GCGACAGTGA CACCAGTGAA GGCAAATGAA 1020 AGTGGAGATT CAGGAGCAGC AATGGTCTCA CCATAGCTGC TGGAATCACA CCTGAGAACT 1080 GAGATATACC AATATTTAAC ATTGTTACAA AGAAGAAAAG ATACAGATTT GGTGAATTTG TTACTGTGAG GTACAGTCAG TACACAGCTG ACTTATGTAG ATTTAAGCTG CTAATATGCT 1200 ACTTAACCAT CTATTAATGC ACCATTAAAG GCTTAGCATT TAAGTAGCAA CATTGCGGTT 1260 TTCAGACACA TOGTGAGGTC CATGGCTCTT GTCATCAGGA TAAGCCTGCA CACCTAGAGT 1320 GTCGGTGAGC TGACCTCACG ATGCTGTCCT CGTGCGATTG CCCTCTCCTG CTGCTGGACT 1380 TCTGCCTTTG TTGGCCTGAT GTGCTGCTGT GATGCTGGTC CTTCATCTTA GGTGTTCATG 1440

	CAGTTCTAAC ACAGTTGGGG TTGGGTCAAT AGTTTCCCAA TTTCAGGATA TTTCGATGTC	1500
	AGAAATAACG CATCTTAGGA ATGACTAAAC AAGATAATGG CAGTTTAGGC TGCACAACTG	1560
5	GTAAAATGAC TGTAGATAAA TGTTGTAATT AGTGTACACG TTTGTATTTT TGTTAATATA	1620
	GCCGCTGCCA TAGTTTTCTA ACTTGAACAG CCATGAATGT TTCATGTCTC CCTTTTTTTT	1680
	TTGTCTATAG CTGTTACCTA TTTTAGTGGT TGAAATGAGA GCTAGTGATG ACAGAAGGAT .	1740
10	GTGGAATGTC TTCTTGACAT CATTGTGTAT TGCTGGTAAT CAAGITGGTA ACGACTACTT	1800
	CTAGCAGCTC TTACCACTAT GACTTAAGTG GTCCTGGAAG GCAGTAAGTG GAGGTTTGCA	1860
15	GCATTCCTGC CTTCATGAGG GCTTCTACCA CTGACCACTT TGCACGTACC TGGCTCCCAG	1920
	ATTTACTTAG GTACCCCACG AGTCGTCCAC ATAAGCAGCT TCATCTTTAC CTTGCCAGAG	1980
20	TIGACAATTA TGGGATACTC TAGTCTACTT ATACTTGTGT TCCCATCTGT CTGCCATCCT	2040
20	CTGAAGGCCA GGACCCAGTC ATACATCCTT AGAAACCAAA GTATGGTTTT TGTTTTCTCT	2100
	TOGAATOTCA GOTCTTAAGG CATTTAATTG AGGGACAAAA AAAAAAAAAA	2160
25	TAGCTAGCTA CTTAAGCATC CATGGGTATT GCTCCATATC AAAGCAGATT TGCAGGACAG	2220
	AAAGAGTAAA TTAGCCTTCA GTCTTGGTTT ACAGCTTCCA AGGAGAGCCT TGGSCACCTG	2280
30	AAATGTTAAC TOGGTCCCTT CCTGTCTCTA GTTCATCAGC ACCTGCAGAT GCCTGACTCT	2340
50	TGTTAGCCTT ACTATTCAAT ACAGTCCTTA GATTCACGGT ATGCCTCTTC CTATCCAGGC	2400
	ACCTATTCTG AATCACCATG TIGCTCTGCA GCTAGAGTTG ATAGGAGAAA ATCCATTTGG	2460
35	GTAGATGGCC TATGAATTTG TAGTAGACTT TCAAAATGAG TGATTTGTTA GCTTGGTACT	2520
	TTTAAGTTTG TGGTACAGAT CCTQCAAACC CATACTCTGA GCAATTAACT GCCTTGAACA	2580
40	TAGAGAAAAA TTAAGGCCTC ACAGGATGAG TCTCCATTCT CTGTAAATGC TTATTTTATC	2640
40	ATAGTOTTA GCCTCTAACT ATGAGTAAAA TGTTCTCTTC GGCCGGGTGT GGTGACTCAC	2700
٠.	ACCTGTAACC TCAGCACTTT GGGAGGCAGA GGTGGGAGGA TCACTTAGGT CCAGGAGTTC	2760
45		282
	GGTGGTATGT ATCTGTGTCC CAGCTAATTG GGAGGGTGAG ATGGGAGGAT TGTTTGAGCC	288
50	TAGGAGAGGG AGGTTGCAGT GAGCCGTGAT CGCACCACTG CACTCCAGCC TGGGCAACAG	294
30	AGCAAGACCC TGTCTTGGAG AAACCAGAAT TTTGGAAGAG CAAATGGGGC TGAGTGCAGT	300
	COMPARED TOTAL	301

60 (i) SEQUENCE CHARACTERISTICS:

⁽²⁾ INFORMATION FOR SEQ ID NO: 221:

	(A) LENGTH: 968 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 221:	
	GGCACGAGGG CCGCGGGACA TCCACGGGGC GCGAGTGACA CGCGGGAGGG AGAGCAGTGT	60
10	TCTGCTGGAG CCGATGCCAA AAACCATGCA TTTCTTATTC AGATTCATTG TTTTCTTTTA	120
	TCTGTGGGCC CTTTTTACTG CTCAGAGACA AAAGAAAGAG GAGAGCACCG AAGAAGTGAA	180
16	AATAGAAGTT TTGCATCGTC CAGAAAACTG CTCTAAGACA AGCAAGAAGG GAGACCTACT	240
15	NAAATGCCCA TTATGACGGC TACCTGGCTA AAGACGGCTC GAAATTCTAC TGCAGCCGGA	300
	CACAAAATGA AGGCCACCCC AAATGGTTTG TTCTTGGTGT TGGGCAAGTC ATAAAAGGCC	360
20	TAGACATTGC TATGACAGAT ATGTGCCCTG GAGAAAAGCG AAAAGTAGTT ATACCCCCTT	420
	CATTTGCATA CGGAAAGGAA GGCTATGCAG AAGGCAAGAT TCCACCGGAT GCTACATTGA	480
25	TTTTTGAGAT TGAACTTTAT GCTGTGACCA AAGGACCACG GAGCATTGAG ACATTTAAAC	540
23	AAATAGACAT GGACAATGAC AGGCAGCTCT CTAAAGCCGA GATAAACCTC TACTTGCAAA	600
	GGGAATTTGA AAAAGATGAG AAGCCACGTG ACAAGTCATA TCAGGATGCA GTTTTAGAAG	660
30	ATATTTTAA GAAGAATGAC CATGATGGTG ATGCCTTCAT TTCTCCCAAG GAATACAATG	720
	TATACCAACA CGATGAACTA TAGCATATTT GTATTTCTAC TTTTTTTTT TAGCTATTTA	780
35	CIGTACTITA TGTATWAAAC AAAGTCMCTT TICTCCMAGT TGKATTTGCT ATTTTTCCCC	840
33	TATGAGAAGA TATTTTGATC TCCCCAATAC ATTGATTTTG GTATAATAAA TGTGAGGCTG	900
	TTTTGCAAAC TTAAAAAAAA ATTTAAAAAA ACTGGAGGG GGCCCGTACC CAANTCGCCG	960
40	NATATGAT	968
45	(2) INFORMATION FOR SEQ ID NO: 222:	
73	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1404 base pairs (B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 222:	
55	CGTTTTCCGG CCGTGCGTTT GTGGCCGTCC GGCCTCCCTG ACATGCAGCC CTCTGGACCC	60
<i></i>	CGAGGITGGA CCCTACTGTG ACACACCTAC CATGCGGACA CTCTTCAACC TCCTCTGGCT	120
	TGCCCTGGCC TGCAGGCCTG TTCACACTAC CCTGTCAAAG TCAGATGCCA AAAAAGCCGC	180
60	IOCCCIOCC IOCHOCCTA LICHTHETHE CETATORNIA ICHINISCON MANNAGOCO	

475

	CTCAAAGACG CTGCTGGAGA AGAGTCAGTT TTCAGATAAG CCGGTGCAAG ACCGGGGTTT	240
	GGTGGTGACG GACCTCAAAG CTGAGAGTGT GGTTCTTGAG CATCGCAGCT ACTGCTCGGC	300
5	AAAGGCCCGG GACAGACACT TTGCTGGGGA TGTACTGGGC TATGTCACTC CATGGAACAG	360
	CCATGGCTAC GATGTCACCA AGGTCTTTGG GAGCAAGTTC ACACAGATCT CACCCGTCTG	420
10	GCTGCAGCTG AAGAGACGTG GCCGTGAGAT GTTTGAGGTC ACGGCCCTCC ACGACGTGGA	. 480
10	CCAAGGGTGG ATGCGAGCTG TCAGGAAGCA TGCCAAGGGC CTGCACATAG TGCCTCGGCT	540
	CCTGTTTGAG GACTGGACTT ACGATGATTT CCGGAACGTC TTAGACAGTG AGGATGAGAT	600
15	AGAGGAGCTG AGCAAGACCG TGGTCCAGGT GGCAAAGAAC CAGCATTTCG ATGGCTTCGT	660
	GGTGGAGGTC TGGAACCAGC TGCTAAGCCA GAAGCGCGTG GGCCTCATCC ACATGCTCAC	720
20	CCACTTGGCC GAGGCTCTGC ACCAGGCCCG GCTGCTGGCC CTCCTGGTCA TCCCGCCTGC	780
20	CATCACCCCC GGGACCGACC AGCTGGGCAT GTTCACGCAC AAGGAGTTTG AGCAGCTGGC	840
	CCCCGTGCTG GATGGTTTCA GCCTCATGAC CTACGACTAC TCTACAGCGC ATCAGCCTGG	900
25	CCCTAATGCA CCCCTGTCCT GGGTTCGAGC CTGCGTCCAG GTCCTGGACC CGAAGTCCAA	960
	GTGGCGAAGC AAAATCCTCC TGGGGCTCAA CTTCTATGGT ATGGACTACG CGACCTCCAA	1020
30	GGATGCCCGT GAGCCTGTTG TCGGGGCCAG GTACATCCAG ACACTGAAGG ACCACAGGCC	1080
30	CCGGATGGTG TGGGACAGCC AGGYCTCAGA GCACTTCTTC GAGTACAAGA AGAGCCGCAG	1140
	TGGGAGGCAC GTCGTCTTCT ACCCAACCCT GAAGTCCCTG CAGGTGCGGC TGGAGCTGGC	1200
35	CCGGGAGCTG GGCGTTGGGG TCTCTATCTG GGAGCTGGCC AGGGCCTGGA CTACTTCTAC	1260
	GACCTGCTCT AGGTGGGCAT TGCGCCTCC GCGGTGGACG TGTTCTTTTC TAAGCCATGG	1320
40	AGTGAGTGAG CAGGTGTGAA ATACAGGCCT NCACTCCGTT TGCTGTGAAA AAAAAAAAAA	1380
40	AAAA AAAAAAAAA AAAA	1404
٠.		
45	(2) INFORMATION FOR SEQ ID NO: 223:	
	(i) SEQUENCE CHARACTERISTICS:	
50	(A) LENGTH: 707 base pairs	
30	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 223:	

NGCGCGCCTG CAGTCGACAC TAGTGGATCC AAAGAATTCG GCACGAGGGC AGGTCCAGGG

CTCAGAAATC AGCTCTATTG ACGAATTCTG CCGCAAGTTC CGCCTGGACT GCCCGCTGGC

CATGGAGGGG ATCAAGGAGG ACCGGCCCAT CACCATCAAG GACGACAAGG GCAACCTCAA

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-	COSCISCATO GCAGACGIGG TOTOGOTOTT CATCACGGIC ACGGACAAGO TGCGCCIGGA	240				
5	GATCCGCGCC ATGGATGAGA TCCAGCCCGA CCTGCGAGAG CTGATGGAGA CCATGCACCG	300				
	CATGAGCCAC CTCCCACCCG ACTITEAGGG CCGCCAGACG GTCAGCTAGT GGCTGCAGAC	360				
	CCTGAGCGGC ATGTCGGCGT CAGATGAGCT GGACGACTCA CAGGTGCGTC AGATGCTGTT	420				
10	CGACCTOGAG TCAGCCTACA ACGCCTTCAA CCGCTTCCTG CATGCCTGAG CCCGGGGCAC	480				
	TAGCCCTTGC ACAGAAGGC AGAGTCTGAG GCGATGGCTC CTGGTCTCTT GTGTGCCACA	540				
15	CAGGCCGTGG TCATCCACAC AACTCACTGT CTGCAGCTGC CTGTCTGGTG TCTGTCTTTG	600				
13	GTGTCAGAAC TTTTGGGCCG GGCCCCTCCC CACAATAAAG ATGCTCTCCG ACCTTCAAAA	660				
	AAAAAAAAA AAAAACTCRG GGGGGGCCCG GTCCCAATCC CCCCM2;	707				
20						
	(2) INFORMATION FOR SEQ ID NO: 224:					
	(2) INFORMATION FOR SEQ ID NO: 224:					
25	(i) SEQUENCE CHARACTERISTICS:					
	(A) LEXCTH: 1334 base pairs					
	(B) TYPE: nucleic acid					
	(C) STFANDEINESS: double (D) TOPOLOGY: linear					
30	(b) foromer. Inter-					
•	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 224:					
	GGGGAACTGC AGTGACAGCA GGAGTAAGAG TUGGAGGCAG GACAGAGTTG GGACACAGGT	60				
35	ATGGAGAGGG GGTTCAGCGA GCCTAGAGAG GGCAGACTAT CAGGGTGTGG GCGGTGAGAA	120				
	TOCAGGGAGA GGAGCGGAAA CAGAAGAGGG GCAGAAGACC GGGGCAUTTG TGGGTTTGCAG	180				
40	ASCCCCTCAG CCATGTTGGG AGCCAAGCCA CACTGGCTAC CAGGTCTTCT ACACAGTCCCC	240				
40	GOOCTGCCCT TOGTTCTGGT GCTTCTGGCC CTGGGGGCCG GGTGGGCTCA GGAGGGTCA	300				
٠,	GASCCCGTCC TGCTGGAGGG GGASTGCCTG GTSGTCTGTG AGCCTGSCCG AGTTGCTGCA	36				
45	COSCOCCO COCCACACO CONTROL DALCOCCOCAC COCCACACOCCO ACTITICATOCCO	42				
	GTCCGAAGCC AMCACCATGA GCCAGCAGGG GAAACCGGCA ATGGCACCAK TGGGGCCATC	48				
50	TACTTCGACC AGGTCCTGGT GAACGAGGGC GGTGGCTTTG ACCGGGCTTC TGGCTCCTTC	54				
JU	GTAGCCCCTG TCCGGGGTGT CTACAGCTTC CGGTTCCATG TGGTGAAGGT GTACAACCGC	60				
	CANACTOTICS AGGIGAGEST GAISSTIGAAS ACGIGGESTS TEATSTICAGE CITTECCAAT	66				
55	GATCCTGACG TGACCCGGGA GGCAGCCACC AGCTCTGTGC TACTGCCTTT GGACCCTGGG	72				

GACCGAGTGT CTCTGCGCCT GCGTCGGGGG AATCTACTGG GTGGTTGGAA ACACTCAAGT

TTCTCTGGCT TCCTCATCTT CCCTCTGA GGACCCAAGT YTTTCAAGCA CAAGAATCCA

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	GCCCCTGACA	ACTITCTICT	GCCCTCTCTT	GCCCCAGAAA	CAGCAGAGGC	AGGAGAGAGA	900
	CTCCCTCTGG	YTCCTATCCC	ACYTCTTTGC	ATGGGAMCCT	GTGCCAAACA	CCCAAGTTTA	960
5	AGARAARARY	ARARCTGWGG	CAGGTATACA	GAGCTGGAAG	TGGACCATGG	AAAACATSGA	1020
	TAACCATGCA	TCYTCTTGCT	TGGCCACCTC	CTGAAACTGT	CCACCTTTGA	AGTTTGAACT	1080
10	TTAGTCCCTC	CAMACTOTGA	CTGCTGCCTC	CTTCCTCCCA	GCTCTCTCAC	TGAGTTATYT	. 1140
10	TCACTGTACC	TGTTCCAGCA	TATCCCCACT	ATCTCTCTTT	CTCCTGATCT	GTGCTGTCTT	1200
	ATTCTCCTCC	TTAGGCTTCC	TATTACCTGG	GATTCCATGA	TTCATTCCTT	CAGACCCTCT	1260
15	CCTGCCAGTA	TGCTAAACCC	TCCCTCTCTC	TTTCTTATCC	CGCTGTCCCA	TTGGCCCAGC	1320
	CTGGATGAAT	CTATCAATAA	AACAACTAGA	GAATGGTGGT	СААААААА	ААААААААА	1380
20	TCGA						1384

(2) INFORMATION FOR SEQ ID NO: 225:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 760 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 225:

GGGTCGACCC ACGCGTCCGC TGACCAGTCC GTTATAGATA CTTCTTCCTA TACCAAAACT 60 GTTTAAACAG GTGCCACCAC AAGGGATGTC GTCCTTACTC TCTGCGGGTC TTCAAGCATC 120 CCTTTGTGGG AAARGTCTCT GGGCAAGCAC GTGGTATTTG GTCTGCTGCT TGCTTCCCTT 180 TTTCCACCAG GGATGTTGTG ATCATAAGTC AAAACAACAG TATATTCCAA ATCTCAAAAG 240 CTATTGTGGC CTGAGCACAA TTGAAATCTA GCAGAGTTTT TCCTATGTAG CTTTAGAGTA 300 ACTOTTOTOC TYCTOTICA CTTACAATTC AGGITCTGCC TYTGCCTAAG AGCATGAGCA 360 GAAGAGTCCT CATGTGACGC TTAGTTCTAT TGCAGTCCTG GGTGAAACTA TTTAAGCWAT 480 GGGCTGCTK CTCCCCANWT CCTCCCTAAC AATTCGTTGT GTGGACTTCT CATCTAAAAG GTTAGTGGCT TTTGCTTGGG ATCAGTGCTC TCTATTGATG TTCTTGCTGG TCTCCAGACA 540 CATTCCTGTT GCATTAAGAC TTGAAAGACT TGTAGATGTG TGATGTTCAG GCACAGGATG 600 CTGAAAGCTA TGTTACTATT CTTAGTTTGT AAATTGTCCT TTTGATACCA TCATCTTGTT 660 720 760 ΑΛΑΛΑΛΑΑ ΑΛΑΛΑΛΑΑ ΝΑΛΑΛΑΛΑΑ ΑΛΑΛΑΛΑΑ

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(2)	INFORMATION	FOR	SEO	ID	NO:	226:

5 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2057 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 226:

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CCGAGCCGCC TGCGCCGGGG GAATCCGTGC GGGCGCCTTC CGTCCCRGTC CCATCCTCGC 60 CGCGCTCCAG CACCTCTGAA GTTTTGCAGC GCCCAGAAAG GAGGCGAGGA ACGAGGGAGT 120 180 AGGGGGGGG CAAAAATGGC TGGGGCAATT ATAGAAAACA TGAGCACCAA GAAGCTGTGC 240 ATTGTTGGTG GGATTCTGCT CGTGTTCCAA ATCATCGCCT TTCTGGTGGG AGGCTTGATT 300 GCTCCAGGGC CCACAACGGC AGTGTCCTAC ATGTCGGTGA AATGTGTGGA TGCCCGTAAG 360 AACCATCACA AGACAAAATG GTTCGTGCCT TGGGGACCCA ATCATTGTGA CAAGATCCGA 420 GACATTGAAG AGGCAATTCC AAGGGAAATT GAAGCCAATG ACATCGTGTT TTCTGTTCAC 480 ATTCCCCTCC CCCACATGGA GATGAGTCCT TGGTTCCAAT TCATGMTGTT TATCCTGCAG 540 CTGGACATTG CCTTCAAGCT AAACAACCAA ATCAGRGAAA ATGCAGAAGT CTCCATGGAC 600 GTTTCCCTGG CTTACCGTGA TGACGCGTTT GCTGAGTGGA CTGAAATGCC CCATGAAAGA 660 GTACCACGGA AACTCAAATG CACCTTCACA TCTCCCAAGA CTCCAGAGCA TGGAGGGCCG 780 GTTACTATGA ATGTGATGTC CTTCCTTTCA TGGAAATTGG GTCTGTGGCC CATGAAGTTT TACCTTTAA ACATCCGGCT GCCTGTGAAT GAGAAGAAGA AAATCAATGT GGGAATTGGG GAGATAAAGG ATATCCGGTT GGTGGGGATC CACCAAAATG GAGGCTTCAC CAAGGTGTGG 900 TTTGCCATGA AGACCTTCCT TACGCCCAGC ATCTTCATCA TTATGGTGTG GTATTGGAGG 960 AGGATCACCA TGATGTCCCG ACCCCCAGTG CTTCTGGAAA AAGTCATCTT TGCCCTTGGG 1020 ATTTCCATGA CCTTTATCAA TATCCCAGTG GAATGGTTTT CCATCGGGTT TGACTGGACC 1080 TOGATGCTGC TGTTTGGTGA CATCCGACAG GCATCTTCTA TGCRATGCTT CTKTCCTTCT 1140 GGATCATCTT CTGTGGGGAG CACATGATGG ATCAGCACGA GCGGAACCAC ATCGCAGGGT 1200 ATTGGAAGCA AGTCGGACCC ATTGCCGTTG GTCCTTCTGC CTCTTCATAT TTGACATGTG 1260 TGAGAGAGGG GTACAACTCA CGAATCCCTT CTACAGTATC TGGACTACAG ACATTGGGAA 1320 CAGAGCTGGC CATGGCTTTC ATCATCGTGG CTGGAATCTG CCTCTGCCTC TAACTTCCTG 1380 TTTCTATGCT TCATGGTATT TCAGGTGTTT CGGAACATCA GTGGGAAGCA GTCCAGCCTG 1440

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	CCASCTATGA	CCYYYLLCC2	GOGGETACAC	TATGAGGGGC	TAATTTTTAG	GTTCAAGTTC	1500
	CTCATGCTTA	TCACCTTGGC	STREEGGTGSS	ATGACTGTCA	TOTTOTTCAT	CGTTAGTCAG	1560
5	GTAACGGAAG	GCCATTGGGA	AATGGGGGGG	COTCACASTO	CCAAGTGAAC	AGTGCCTTTT	1620
	TCACAGGCAT	CTATGGGATG	TOGALITOTOT	AUGICITIEC	TOTGATGTTC	TTGTATGCAC	1680
10	CATCCCATAA	AAACTATGGA	GAAGACCAST	CCAATGGAAT	GCAACTCCCA	TGTAAATCGA	1740
10	GGGAAGATTG	recessorer	GTTTCGGAAC	TTTATCAAGA	ATTGTTCAGC	GCTTCGAAAT	1800
	ATTCCTTCAT	CAATGACAAD	SCHOOLLACE	GIATTIGAGE	CAACAAGGCA	ACACATGTTT	1860
15	ATCAGCTTTG	CATTIGOAGT	TATCHENETO	ACATTGATTG	TACTTGTATA	CGCACACAAA	1920
	TACACTCATT	TAGCCTTTAT	CLCYYYYCCL	TANATATAAS	GAAAAAAGCG	TCAACAATAA	1980
20	ATATICTITG	AGUATIGICI	TACTFCTCTT	AAAAAAAAA	AAAAAAACTC	GTGCCGAATT	2040
20	CGGCACGAGG	GGCACGA					2057

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(2) DEFORMATION FOR SEC ID NO: 227:

(i) SEQUENCE THARACTERISTICS:

(A) LENGTH: 2084 base pairs

(B) TYPE: nucleic acid

(C) STRAICEMMESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 227:

35 GOCAGAGGGG CANTIFOCTIGG AFAGAGGGAA ACCCCCATTC CTCTGTGCCC CTCCTGTCCC 60 ACCARGIGCT TERTHARART AUCTOTTGIT ACCGGRARATA ACTGITCATT TITCACTCCT 120 CCCTCCTAGG TCACACTTTT CAGAAAAAAA ATCTGCATCC TGGAAACCAG AAGAAAAATA 40 180 TORGROSCOG REMONICOUR TERRETETET SCIENCIFFE GOTGROTTETE TOGRETCOTO 240 CTCAGGTGTT AGGTACAGTG TGTTTGATCG TGGTGGCTTG AGGGGAACCG CTTGTTCAGA 45 GCTGTGACTG CGCCTGCACT GCAGAGAAGC TGCCCTTTGGC TGCTCGTAGC GCCGGGCCTT 360 CTCTCCTCGT CRTCATCCAG ASCAGCCAGT GTCCGGGAGG CAGAAGGTAC CGGGGCAGCT 50 ACTGGLEGAL TGTGCGGGCC TGCCTGGGCT GCCCCCTCCG CCGTGGGGGCC CTGTTGCTGC 480 TGTCCATCTA TETCTACTAC TSCCTCCCAA ATGCGGTCGG CCCGCCCTTC ACTTGGATGC 540 TIGCCCTCCT GGGCCTTCTC GCAGGCACTG AACATCCTCC TGGGCCTCAA GGGCCTGGCC 600 55 CCAGCTGAGA TCTCTGCAGT GTGTGAAAAA GGGAATTTCA ACGTGGCCCA TGGGCTGGCA 660 TOGTCATATT ACATCSGACA TOTGCCGCTG ATCOTGCCAG AGCTCCAGGC CCGGATTCGA 720 780 ACTTACAATT AGCATTACAA CAACCTGCTA CGGGGTGCAG TGAGCCAGCG GTGTNATATT 60

480

	CTCCTCCCAT TGGACTGTGG GGTGCCTGAT AACCTGAGTA TGGCTGACCC CAACATTCGC	840
_	TTCCTGGATA AACTGCCCCA GCAGACCGGT GACCGTGCTG GCATCAAGGA TCGGGTTTAC	900
5	AGCAACAGCA TCTATGAGCT TCTGGAGAAC GGGCAGCGGG CGGGCACCTG TGTCCTGGAG	960
	TACGCCACCC CCTTGCAGAC TTTGTTTGCC ATGTCACAAT ACAGTCAAGC TGGCTTTAGC	. 1020
10	GGGGAGGATA GGCTTGAGCA GGCCAAACTC TTCTGCCGGA CACTTGAGGA CATCCTGGCA	1080
	GATGCCCCTG AGTCTCAGAA CAACTGCCGC CTCATTGCCT ACCAGGAACC TGCAGATGAC	1140
1.5	AGCAGCTTCT CGCTGTCCCA GGAGGTTCTC CGGCACCTGC GGCAGGAGGA AAAGGAAGAG	1200
15	GTTACTGTGG GCAGCTTGAA GACCTCAGCG GTGCCCAGTA CCTCCACGAT GTCCCAAGAG	1260
	CCTGAGCTCC TCATCAGTGG AATGGAAAAG CCCCTCCCTC TCCGCACGGA TTTCTCTTGA	1320
20	GACCCAGGGT CACCAGGCCA GAGCCTCCAG TGGTCTCCAA GCCTCTGGAC TGGGGGCTCT	1380
	CTTCAGTGGC TGAATGTCCA GCAGAGCTAT TTCCTTCCAC AGGGGGCCCTT GCAGGGAAGG	1440
25	GTCCAGGACT TGACATCTTA AGATGCGTCT TGTCCCCTTG GGCCAGTCAT TTCCCCTCTC	1500
25	TGAGCCTCGG TGTCTTCAAC CTGTGAAATG GGATCATAAT CACTGCCTTA CCTCCCTCAC	1560
	GGTTGTTGTG AGGACTGAGT GTGTGGAAGT TTTTCATAAA CTTTGGATGC TAGTGTACTT	1620
30	AGGGGGTGTG CCAGGTGTCT TTCATGGGGC CTTCCAGACC CACTCCCCAC CCTTCTCCCC	1680
	TTCCTTTGCC CGGGGACGCC GAACTCTCTC AATGGTATCA ACAGGCTCCT TCGCCCTCTG	1740
25	GCTCCTGGTC ATGTTCCATT ATTGGGGAGC CCCAGCAGAA GAATGGAGAG GAGGAGGAGG	1800
35	CTGAGTTTGG GGTATTGAAT CCCCCGGCTC CCACCCTGCA GCATCAAGGT TGCTATGGAC	1860
	TCTCCTGCCG GGCAACTCTT GCGTAATCAT GACTATCTCT AGGATTCTGG CACCACTTCC	1920
40	TTCCCTGGCC CCTTAAGCCT AGCTGTGTAT CGGCACCCCC ACCCCACTAG AGTACTCCCT	1980
	CTCACTTGCG GTTTCCTTAT ACTCCACCCC TTTCTCAACG GTCCTTTTTT AAAGCACATC	2040
45	TCAGATTAAA AAAAAAAAA AAAAAAAAAA AGGGGGGCN GCNT	2084

(2) INFORMATION FOR SEQ ID NO: 228:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2143 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 228:

TOGACCCACG CGTCCGGTTG AATTCCTTGA CCTGCAAACA CATATTTATT AGCCTGACTC

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•	AAACAATGAA	GCTATTAAAA	CTTCGGAGGA	ACATTGTAAA	ACTOTOTTTG	TATCGGCATT	120
	TCACCAACAC	GCTTATTTTG	GCAGTGGCAG	CATCCATTGT	GTTTATCATC	TGGACAACCA	180
5	TGAAGTTCAG	AATAGTGACA	TGTCAGTCGG	ACTGGCGGGA	GCTGTGGGTA	GACGATGCCA	240
	TCTGGCGCTT	GCTGTTCTCC	ATGATCCTCT	TIGICATCAT	GGTTCTCTGG	CGACCATCTG	300
10	CAAACAACCA	GAGGTTTGCC	TTTTCACCAT	TGTCTGAGGA	AGAGGAGGAG	GATGAACAAA	360
10	AGGAGCCTAT	GCTGAAAGAA	AGCTTTGAAG	GAATGAAAAT	GAGAAGTACC	AAACAAGAAC	420
	CCAATGGAAA	TAGTAAAGTT	AACAAAGCAC	AGGAAGATGA	TTTGAAGTGG	GTAGAAGAGA	480
15	ATGTTCCTTC	TTCTGTGACA	GATGTAGCAC	TTCCAGCCCT	TCTGGATTCA	GATGAGGAAC	540
	GAATGATCAC	ACACTTTGAA	AGGTCCAAAA	TGGAGTAAGG	AATGGGAAGA	TTTGCAGTTA	600
20	AAGATGGCTA	CCATCAGGGA	AGAGATCAGC	ATCTGTGTCA	GTCTTCTGTA	CGGCTCCATG	660
20	GGATTAAAGG	AAGCAATGAC	ATCCTGATCT	GTTCCTTGAT	CTTTGGGCAT	TGGAGTTGGC	720
	GAGAGGTGTC	AGAACAAAGA	GAACATCTTA	CTGAAAACAA	GTTCATAAGA	TGAGAAAAAT	780
25	CTACGAGCTT	CTTATTTACA	ACACTGCTGC	CCCCTTTCCT	CCCAGACTCT	GACATGGATG	840
	TTCATGCAAC	TTAAGTGTGT	TGTTCCTGAA	CTTTCTGTAA	TGTTTCATTT	TTTAAATCTG	900
30	ACAAACTAAA	AAGTTTAACG	TCTTCTAAAA	GATTGTCATC	AACACCATAA	TATGTAATCT	960
50	CCAGGAGCAA	CTGCCTGTAA	TTTTTTTTA	TTTAGGGAGT	TACATAGGTG	ATGGGGGAAA	1020
	TTGTTAACTA	CCTTTCATTT	TCCTGGGAAG	TCAAGGTTAC	ATCTTGCAGA	CCTTCTTTTC	1080
35	AGAAAAAAGG	GCCCTTCTGA	GTTAAGGAGC	CATAGITCTA	TCAATGATCA	AAAGAAAAA	1140
	AAAAAAAAGA	GAAACTGTTA	CAGTATGATT	CAGATCATTT	AAAAAAGCAA	AATCAAGTGC	1200
40	AATTTTGTTT	ACAAATGGTG	TATATTAAAG	ATTTTTCTAT	TTCAGATGTA	CTTTAAAGAG	1260
. ,	AAATATTAGC	TTAACTCTTT	TGACATCTGC	TATTGTGACA	CATCCCATTG	CTGGCAATGT	1320
	GGTGCACACT	CCGAAACTTT	TAACTACTGT	TTTGTAAGCC	TCCAAGGGTG	GCATTGCAGG	1380
45	GTCCTTAGGC	AATGTTTTGT	TTGCCTTTAT	GCAGAGAGGT	GCTCCAAGTG	CTGTGATTGA	1440
	GCACCGTGCT	AGAGGAACTG	TAATGCTTCA	GAAGTIGTAG	CTTATACAAA	GGAAACAGGT	1500
50	CCTGCTGGCT	TAATTTAAAC	AGITATTGCA	TGAAGTAGCG	TGGAGGCCCT	GGACTGCTGC	1560
	TCGTTCTTTA	GGATGGACTG	TTCTGGTATC	TGGTATTGGT	TTAGAGACTG	TTAATAAGGG	1620
	ACATCACAAG	GTGATGGGAT	TCATTTGAAG	CACTCTATTT	CTGTTTTAAT	GGTTTTATCC	1680
55	AATTTTGCCT	TCCCAAGATT	TTTGTTCTAC	ATAAAAAGTT	CATGCCACTT	TITAATATAA	1740
	AAAATTTAA	CAAAATTAAT	GTATTTTCT	CATTTTTTC	AAACTTTTTC	TAAAGACTCT	1800
60	TTCTGTCAAA	CTCATGAAAA	ATTTCTTTCT	ATGGCTTTTA	TTCTAGATTG	TCTTATTTTC	1860

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TGTTAAAACC	AATGACCACA	TGACCACAAT	CTTCACTAAC	TCATACTGCA	GTGAAAGTGT	1920
TAACCCTTAG	GTAGTTTCTC	TACAACTCTT	TGCTATGGTG	AAAATTTTTA	AAGTTTCCTA	1980
GGGAAGTATC	TCTGAGGGAA	CAGGCAATCT	GAAGGAACTG	ACTATATTCT	CCATGGCTAA	2040
GTCCATTAGG	CCAAAAGNCT	GGGTGGGTAT	TOGTTGTCAN	GCTGTCTATT	GGCATATTAA	2100
AAACGTAGGC	CGGANGGAAT	AATTAGGTTG	TNATGCCGGC	GGG		2143

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(2) INFORMATION FOR SEQ ID NO: 229:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1025 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double

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(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 229:

25	CCTGGCCCAC ATTGCTTCAT TGGCCTGGCC ATGCGCCTGT ACTATGGCAG CCGCTAGTCC	60
	CTGACAACTT CCACCCTGAT TCCGGACCCT GTAGATTGGG CGCCACCACC AGATCCCCCT	120
	CCCAGGCCTT CCTCCCTCTC CCATCAGCAG CCCTGTAACA AGTGCCTTGT GAGAAAAGCT	180
30	GGAGAAGTGA GGGCAGCCAG GTTATTCTCT GGAGGTTGGT GGATGAAGGG GTACCCTAGG	240
	AGATGTGAAG TGTGGGTTTG GTTAAGGAAA TGCTTACCAT CCCCCACCCC CAACCAAGTT	300
25	CTTCCAGACT AAAGAATTAA GGTAACATCA ATACCTAGGC CTGAGAAATA ACCCCATCCT	360
35	TGTTGGGCAG CTCCCTGCTT TGTCCTGCAT GAACAGAGTT GATGAAAGTG GGGTGTGGCC	420
	AACAAGTGGC TTTCCTTGCC TACTTTAGTC ACCCAGCAGA GCCACTGGAG CTGGCTAGTC	480
40	CAGCCCAGCC ATGGTGCATG ACTCTTCCAT AAGGGATCCT CACCCTTCCA CTTTCATGCA	540
٠.	AGAAGGCCCA GTTGCCACAG ATTATACAAC CATTACCCAA ACCACTCTGA CAGTCTCCTC	600
45	CAGTTCCAGC AATGCCTAGA GACATGCTCC CTGCCCTCTC CACAGTGCTG CTCCCCACAC	660
43	CTAGCCTTTG TTCTGGAAAC CCCAGAGAGG GCTGGGCTTG ACTCATCTCA GGGAATGTAG	720
•	CCCCTGGGCC CTGGCTTAAG CCGACACTCC TGACCTCTCT GTTCACCCTG AGGGCTGTCT	780
50	TGAAGCCCGC TACCCACTCT GAGGCTCCTA GGAGGTACCA TGCTTCCCAC TCTGGGGCCT	840
	GCCCCTGCCT AGCAGTCTCC CAGCTCCCAA CAGCCTGGGG AAGCTCTGCA CAGAGTGACC	900
55	TGAGACCAGG TACAGGAAAC CTGTAGCTCA ATCAGTGTCT CTTTAACTGC ATAAGCAATA	960
55	AGATCTTAAT AAAGTCTTCT AGGCTGTAGG GTGGTTCCTA CAACCACAGC CAAAAAAAAA	1020
	AAAA	1025

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(2)	TNFORMATION	FOR	SEO	ID	NO:	230:

5	(i) SEQUENCE CHARACTERISTICS:
_	(A) LENGTH: 1250 base pairs
	(B) TYPE: nucleic acid
	(C) STRANDEDNESS: double
	(D) TOPOLOGY: linear
10	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 230:

GCCCACGCGT CCGCCCACGC GTCCCGCGGT GCGGAGTATG GGGCGCTGAT GGCCATGGAG 60 GGCTACTGGC GCTTCCTGGC GCYGCTGGGG TCGGCACTGC TCGTCGGCTT CCTGTCGGTG 120 ATSTTCGCCC TCGTCTGGGT CCTCCACTAC CGAGAGGGGC TTGGCTGGGA TGGGAGCGCA 180 CTAGAGTTTA ACTGGCACCC AGTGCTSATG GTCACCGGCT TCGTCTTCAT CCAGGGCATC 240 GCATCATCGT CTACAGACTG CCGTGGACCT GGAAATGCAG CAAGCTCCTG ATGAAATCCA 300 TOCATGCAGG GITAAATGCA GITGCTGCCA TICTTGCAAT TATCTCTGTG GIGGCCGTGT 360 TTGAGAACCA CAATGTTAAC AATATAGCCA ATATGTACAG TCTGCACAGC TGGGTTGGAC 420 TGATAGCTGT CATATGCTAT TTGTTACAGC TTCTTTCAGG TTTTTCAGTC TTTCTGCTTC CATGGGCTCC GCTTTCTCTC CGAGCATTTC TCATGCCCAT ACATGTTTAT TCTGGAATTG 540 TCATCTTTGG AACAGTGATT GCAACAGCAC TTATGGGATT GACAGAGAAA CTGATTTTTT 600 CCCTGAGAGA TCCTGCATAC AGTACATTCC CGCCAGAAGG TGTTTTCGTA AATACGCTTG 660 GCCTTCTGAT CCTGGTGTTC GGGGCCCTCA TTTTTTGGAT AGTCACCAGA CCGCAATGGA 720 780 AACGTCCTAA GGAGCCAAAT TCTACCATTC TTCATCCAAA TGGAGGCACT GAACAGGGAG CAAGAGGTTC CATGCCAGCC TACTCTGGCA ACAACATGGA CAAATCAGAT TCAGAGTTAA 840 ACARTGAAGT AGCAGCAAGG AAAAGAAACT TAGCTCTGGA TGAGGCTGGG CAGAGATCTA 900 CCATGTAAAA TGTTGTAGAG ATAGAGCCAT ATAACGTCAC GTTTCAAAAC TAGCTCTACA 960 GTTTTGCTTC TCCTATTAGC CATATGATAA TTGGGCTATG TAGTATCAAT ATTTACTTTA 1020 ATCACAAAGG ATGGTTTCTT GAAATAATTT GTATTGATTG AGGCCTATGA ACTGACCTGA 1080 ATTGGAAAGG ATGTGATTAA TATAAATAAT AGCAGATATA AATTGTGGTT ATGTTACCTT 1140 TATCTTGTTG AGGACCACAA CATTAGCACG GTGCCTTGTG CAKAATAGAT ACTCAATATG 1200 1250 TGAATATGTG TCTACTAGTA GITAATTGGA TAAACTGGCA GCATCCCTGA

(2) INFORMATION FOR SEQ ID NO: 231:

(i) SEQUENCE CHARACTERISTICS:

484

(A) LENGTH: 1811 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 231:

	CNGNCAGTAC CGGTCNGATT CCCGGGTCGA CCCACGCGTC CGCTGCATTC CAGGGCCTTT	60
10	CAGTGGCTTT CATTCTGAAG TTCCTGGATA ACATGTTCCA TGTCTTGATG GCCCAGGTTA	120
	CCASTGTCAT TATCACAACA GTGTCTGTCC TGGTCTTTGA CTTCAGGCCC TCCCTGGAAT	180
	TTTTCTTGGA AGCCSCATCA GTCSTYCTCT CTATATTTAT TTATAATGCC AGCAAGCCTC	240
15	AAGTTCCGGA ATACGCACCT AGGCAAGAAA GGATCCGAGA TCTAAGTGGC AATCTTTGGG	300
	ACCOTTCCAG TGGGGATGGA GAAGAACTAG AAAGACTTAC CAAACCCAAG AGTGATGAGT	360
20	CAGATGAAGA TACTITCTAA CIGGTACCCA CATAGTITGC AGCTCTCTIG AACCTTATIT	420
	TCACATTITC AGIGITIGIA ATATTTATCT TTTCACTTIG ATAAACCAGA AATGITTCTA	480
25	AATCCTAATA TTCTTTGCAT ATATCTAGCT ACTCCCTAAA TGGTTCCATC CAAGGCTTAG	540
25	AGTACCCAAA GGCTAAGAAA TTCTAAAGAA CTGATACAGG AGTAACAATA TGAAGAATTC	600
	ATTAATATCT CAGTACTTGA TAAATCAGAA AGITATATGT GCAGATTATT TTCCTTGGCC	660
30	TTCAAGCTTC CAAAAAACTT GTAATAATCA TGTTAGCTAT AGCTTGTATA TACACATAGA	720
	GATCAATTIG CCAAATATIC ACAATCATGI AGITCIAGIT TACATGCCAA AGICITCCCT	780
25	TTTTAACATT ATAAAAGCTA GGTTGTCTCT TGAATTTTGA GGCCCTAGAG ATAGTCATTT	840
35	TGCAAGTAAA GAGCAACGGG ACCCTTTCTA AAAACGTTGG TTGAAGGACC TAAATACCTG	900
	GCCATACCAT AGATTTGGGA TGATGTAGTC TGTGCTAAAT ATTTTGCTGA AGAAGCAGTT	960
40	TCTCAGACAC AACATCTCAG AATTTTAATT TTTAGAAATT CATGGGAAAT TGGATTTTTG	1020
٠.	TAATAATCTT TIGATGTTTT AAACATTGGT TCCCTAGTCA CCATAGTTAC CACTTGTATT	1080
4.5	TTAAGTCATT TAAACAAGCC ACGGTGGGGC TTTTTTCTCC TCAGTTTGAG GAGAAAAATC	1140
45	TTGATGTCAT TACTCCTGAA TTATTACATT TTGGAGAATA AGAGGGCATT TTATTTTATT	1200
	AGITACIAAT TCAAGCTGTG ACTATTGTAT ATCTTTCCAA GAGTTGAAAT GCTGGCTTCA	1260
50	GAATCATACC AGATTGTCAG TGAAGCTGAT GCCTAGGAAC TTTTAAAGGG ATCCTTTCAA	1320
	AAGGATCACT TAGCAAACAC ATGTTGACTT TTAACTGATG TATGAATATT AATACTCTAA	1380
ـ ـ	ARATAGARAG ACCAGTARTA TATRAGTCAC TITACAGTGC TACTTCACAC TTARARGTGC	1440
55	ATGGTATTT TCATGGTATT TTGCATGCAG CCAGTTAACT CTCGTAGATA GAGAAGTCAG	1500
	GTGATAGATG ATATTAAAAA TTAGCAAACA AAAGTGACTT GCTCAGGGTC ATGCAGCTGG	1560
60	THE PARTY OF THE P	1620

485

•	TARATATGAG CITTATGGTG TCATTCTCAG ARACTTATAC ATTTCTGCTC TCCTTTCTCC	1680
_	TAAGTTTCAT GCAGATGAAT ATAAGGTAAT ATACTATTAT ATAATTCATT TGTGATATCC	1740
5	ACAATAATAT GACTGGCAAG AATTGGTGGA AATTTGTAAT TAAAATAATT ATTAAACCTA	1800
	AAAAAAAAN N	. 1811
10		
	222	
	(2) INFORMATION FOR SEQ ID NO: 232:	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2271 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 232:	
	CTGACCTCAT GGCGTAGAGC CTAGCAACAG CGCAGGCTCC CAGCCGAGTC CGTTATGGCC	60
25	GCTGCCGTCC CGAAGAGGAT GAGGGGCCA GCACAAGCGA AACTGCTGCC CGGGTCGGCC	120
	ATCCAAGCCC TIGIGGGGTT GGCGCGGCCG CTGGTCTTGG CGCTCCTGCT TGTGTCCGCC	180
20	GCTCTATCCA GTGTTGTATC ACGGACTGAT TCACCGAGCC CAACCGTACT CAACTCACAT	240
30	ATTICTACCC CAAATGIGAA TGCTTTAACA CATGAAAACC AAACCAAACC TTCTATTTCC	300
	CAAATCAGCA CCACCCTCCC TCCCACGACG AGTACCAAGA AAAGTGGAGG AGCATCTGTG	360
35	GTCCCTCATC CCTCGCCTAC TCCTCTGTCT CAAGAGGAAG CTGATAACAA TGAAGATCCT	420
	AGTATAGAGG AGGAGGATCT TCTGATGCTG AACAGTTCTC CATCCACAGC CAAAGACACT	480
40	CTAGACAATG GCGATTATGG AGAACCAGAC TATGACTGGA CCACGGGCCC CAGGGACGAC	540
	GACGAGTCTG ATMGACACCT TOGAAGAAAA CAGGGGTTAC ATGGAAATTG AACAGTCAGT	600
	GAAATCTTTT AAGATGCCAT CCTCAAATAT AGAAGAGGAA GACAGCCATT TCTTTTTTCA	660
45		720
	GATTYTTCTT CTGGTTCAAA GCAGGAAATG GCGTGATGGC CTTTGTTCCA AAACAGTGGA	780
50	ATACCATCGC CTAGATCAGA ATGITAATGA GGCAATGCCT TCTTTGAAGA TTACCAATGA	840
50	TTATATTTT TAAAGCACTG TGATTTGAAT TTGCTTATGT AATTTTATTT GCTTGACTTT	900
	TTATATGATA TTGTGCAAAT GTTTGCCATA GGCAATTGGT ACTTAAATGA GAGGTGAGTC	960
55		
	TGTACTTTTA GAGCTGAGTT TAATCAGGTG TCCAAAATGT GAGTTAAACA TTACCTTATA	1080

TITACACTGT TAGITTITAT TGTTTTAGAT TTATTATGCT TCTTCTGGAA GTATTAGTGA

60

300

	TGCTACTTTT AAAAGATCCC AAACTTGTAA CTAAATTCTG ACATATCTGT TACTGCTGAC	1200
	TCACATTCAT TCTCCGCCAT TCAAATACTA TTTTTTATCC ACATTTTTT TTGTTCCCAA	1260
5	ACTGTAATGT ACAAGGATAT GTGTGATAAT GCTTTGGATT TGAGTAATAT TTTTTTTTCT	1320
	TCCAAGAAAA CTGCTTTGGA TATTTTTAGA TAATTTAAAC ATAATTTAGG ATAATGATAT	1380
	TGCTCAATCT GACCACAATT TTAGGTAAAA CATTAAATGT GTCAAGAAAT CTTGGCAACA	1440
.0	GAGACTICTICC AGCTTGCAGT GGACATAGAT AAAATGTTAC AGAGATACTA TTTTTTTGGT	1500
	TGGAATTACT ATATTAAATT TAGAAGCAGA AACTGGTAAA ATGTTAAATA CATGTACAAT	1560
15	TGCTTTTAGT TAGCAATTGA TTGTAGCATG GGTTCCTCCA AGGTTTCAAG CAATGGGCAG	1620
	AGTTTAAAAT TATATCAGAT TCGTTTACTT CGTTTATTAT TTTACAGTAA ATTTGAATAA	1680
	ATCTTAGGG TCATTATCAC TTAAATAATA CTGTACCTAG GTCTTTCAAA TTAAAATTAT	1740
20	ACCTGAATGA AGTTGTTTGT ATACATAAAG GATATTTGTG TACAATTACC TTTTTTCCCC	1800
	CACACTTGTT TTCTTTGTTT TTGTTTTTTA TGGCAACTGG AAAGTATTTA CTATGGGATT	1860
25	CATTTATGTC TGTCTTCTA TCATAAAGAA TIGATCAATA TGTAAATATG TGATTTGAAC	1920
	CATGGITGAC TTACAAGTGT CACTACAGCT TTTTAGAAAA CATAGCCCTA ATATATGTTA	1980
	AGCAGGACCC GGGTGAGCCA GTGGGCTTGC GCTTTATGTA GAGCTGGAAG AAGGCCGTCC	2040
30	ATCCTGTCTC TTGGGCGGAC AGTGTACTTT CCTAATAGGG AAGGGAAGCA CAATGGAAAT	2100
	ACCCCTGAAC CGTTTTATTG CAGTAATTTT TTTCATATCT GAAACTATTA TTTAATATTT	2160
35	ТСВАТАВСЯТ ТІТАВАВАЯТ ВАВТОССАВА СЯТАТАВАТС ТАВАВАВАВА ВАВАВАВАВА	2220
	и апапабава бабабаба бабабаба бабабаба бабабаба	2271
	·	
40		
	(2) INFORMATION FOR SEQ ID NO: 233:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1338 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 233:	
	CTTCCGGTTC TCCGGGCAGC TGCCACTGCT GTAGCTTCTG CCACCTGCCA CGACCGGGCC	60
	TCTCCCTGGC GTTTGGTCAC CTCTGCTTCA TTCTCCACCG CGCCTATGGT CCCTCTTGGA	120
55	CCCACCTICG CCNCCCTICGC GCCTCCCCGC TGGTGAGAGA GCGGTCCGGG AACGATGAAG	180
	SCCTCSCAST SCTSCTSCTG TCTCASCCAC CTCTTGSCTT CCGTCCTCCT CCTCCTGTTG	240

CTGCCTGAAC TAAGCGGGYC CCTGGMAGTC CTGCTGCAGG CAGCCGAGGC CGCGCCAGGT

•	YTTGGGCCTC CTGACCCTAG ACCAGGACAT TACCGCCGCT GCCACCGGGC CCTWACCCCT	360
5	GCCCAGCAGC CGGGCCGTGG TCTGGCTGAA GCTGCGGGGG CCGCGGGGCT CCGAGGGAGG	420
	CAATGGCAGC AACCCTGTGG CCGGGCTTGA GACGGACGAT CACGGAGGGA AGGCCGGGGA	480
	ARGCTCGGTG GGTGGCGGCC TTGCTGTGAG CCCCAACCCT GGCGACAAGC CCATGACCCA	. 540
10	GOGGGCCCTG ACCGTGTTGA TGGTGGTGAG CGGCGCGGTG CTGGTGTACT TCGTGGTCAG	600
	GACCGTCAGG ATGAGAAGAA GAAACCGAAA GACTAGGAGA TATGGAGTTT TGGACACTAA	660
15	CATAGAAAAT ATGGAATTGA CACCTTTAGA ACAGGATGAT GAGGATGATG ACAACACGTT	720
13	GTTTGATGCC AATCATCCTC GAAGATAAGA ATGTGCCTTT TGATGAAAGA ACTTTATCTT	780
	TCTACAATGA AGAGTGGAAT TTCTATGTTT AAGGAATAAG AAGCCACTAT ATCAATGTTG	840
20	GGGGGTATT TAAGTTACAT ATATTINAAC AACCTTTAAT TTGCTGTTGC AATAAATACC	900
	GTATCCTTTT ATTATATCTT TATATGTATA GAAGTACTCT GTTAATGGGC TCAGAGATGT	960
25	TGGGGATAAA GTATACTGTA ATAATTTATC TGTTTGAAAA TTACTATAAA ACGGTGTTTT	1020
23	CTGRTCGGTT TTTGTTTCCT GCTTACCATA TGATTGTAAA TTGTTTTATG TATTAATCAG	1080
	TTAATGCTAA TTATTTTTGC TGATGTCATA TGTTAAAGAG CTATAAATTC CAACAACCAA	1140
30	CTGGTGTGTA AAAATAATTT AAAATYTCCT TTACTGAAAG GTATTTCCCA TTTTTGTGGG	1200
	GAAAAGAAGC CAAATTTATT ACTITGTGTT GGGGTTTTTA AAATATTAAG AAATGTCTAA	1260
35	GITATIGITI GCAAAACAAT AAATAIGAIT TIAAAITCIC TIAAAAAAAA AAAAAAAAAC	1320
33	CCCGGGGGG GGCCCGGN	1338
40	(2) INFORMATION FOR SEQ ID NO: 234:	
٠.	(i) SEQUENCE CHARACTERISTICS:	
45	(A) LENGTH: 31 amino acids (B) TYPE: amino acid	
73	(D) TOPOLOGY: linear	
•	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 234:	
	Met Leu Ser Thr Gly Ile Glu Val Ala Arg Pro Pro Ala Thr Leu Leu	

Met Leu Ser Thr Gly Ile Glu Val Ala Arg Pro Pro Ala Thr Leu Leu
50 1 5 10 15

Gly Leu Met Phe Val Leu Thr Gly Met Pro Arg Gly Leu Arg Xaa 20 25 30

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(2) INFORMATION FOR SEQ ID NO: 235:

(i) SEQUENCE CHARACTERISTICS:

60

(A) LENGTH: 116 amino acids

	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 235:
5	Met Asn Val Val Ile Val Ile Ile Leu Phe Ser Phe Asp Ser Val Gly 1 5 10 15
	Thr Met Phe Ser Cys Asn Arg Ile Pro Lys Ile Thr Val Leu Asn Lys 20 25 30
10	Leu Lys Phe Xaa Cys Glu Val Leu Leu Arg Ile Gln Thr Ile Gln Gly 35 40 45
15	Phe Tyr Arg Cys Thr Arg Ile Ser Arg Tyr Lys Gly Ile Phe Pro Asp 50 55 60
	Phe Cys Gln Ser Gln Cys Met Gly Cys Asn Pro Glu Ser Xaa Met Ala 65 70 75 80
20	Val Pro Ala Leu Val Thr Pro Ile Leu Ala His Arg Lys Lys Glu Lys 85 90 95
25	Gly Met Cys Leu Phe Thr Leu Ile Ile Ala Pro Thr Arg Cys Thr His 100 105 110
25	Tyr Phe Cys Xaa 115
30	(2) INFORMATION FOR SEQ ID NO: 236:
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 103 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 236:
40	Met Ser Ser Ala Lys Ile Val Arg Gln Arg Gly Ala Val Pro Thr Tyr 1 5 10 15
٠.	Tyr Thr Thr Glu Ala Gly Glu Ile Ile Phe Leu Val Leu Asn Trp Ser 20 25 30
45	Leu Ser Ile Leu His Ile Val Asp Val Leu Cys Ser Lys Pro Glu Lys 35 40 45
	Ser Val Thr Glu Asp Ala Ala Ser Gly Leu Ser Gln Arg Met Thr Ala 50 55 60
50	Leu Val Trp Arg Lys Gly Pro Asp Gly Gly Ser Arg Lys Pro Ile Le 65 70 75 8
55	Leu Leu Phe Phe Phe Leu Pro Leu Ile Leu Cys Phe His Ser Phe Il 85 90 95
	His Ser Ser Asn Ile Cys Xaa 100

	(2) INFORMATION FOR SEQ ID NO: 237:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 42 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 237:
10	Met Ile Leu Phe Pro Gln Xaa Ala Leu Arg Leu Gly Xaa Trp Pro Arg 1 5 10 15
15	Thr Trp Ser Ile Leu Xaa Lys Tyr Ser Val Asn Phe Phe Ser Ala Tyr 20 25 30
••	Ser Pro Met Gly Ala Val Gly Thr Glu Phe 35 40
20	(2) INFORMATION FOR SEQ ID NO: 238:
25	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 37 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 238:
30	Met Ile Ile Leu Leu Leu Phe Met Leu Leu Asn Asn Val Val Leu Val 1 5 10 15
	Gln Glu Asp Asn Cys Gln Arg Lys Asn Thr Val Gln Glu Arg Arg Xaa 20 25 30
35	Trp Ser Gln Trp Xaa 35
40	(2) INFORMATION FOR SEQ ID NO: 239:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 128 amino acids
	(B) TYPE: amino acid
45	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 239:
50	Met Ala Ala Xaa Pro Pro Gly Cys Thr Pro Pro Xaa Leu Leu Asp Ile 1 5 10 15
50	Ser Trp Leu Thr Glu Ser Leu Gly Ala Gly Gln Pro Val Pro Val Glu 20 25 30
55	Cys Arg His Arg Leu Glu Val Ala Gly Pro Arg Lys Gly Pro Leu Ser 35 40 45
	Pro Ala Trp Met Pro Ala Tyr Ala Cys Gln Arg Pro Thr Pro Leu Thr 50 55 60
60	His His Asn Thr Gly Leu Ser Glu Leu Leu Glu His Gly Val Cys Glu

	Glu Val Glu Arg Val Arg Arg Ser Glu Arg Tyr Gln Thr Met Lys Val 85 90 95
5	Arg Arg Ala Gly Leu Gly Pro Thr Pro Gly Met Ser Cys Pro Gly Asn 100 105 110
10	Asp Asn Thr Val His Thr Met His Gly Glu Ala Asn Arg Gly Ser Xaa 115 120 125
15	(2) INFORMATION FOR SEQ ID NO: 240:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 67 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 240:
25	Met Ser Ile Leu Cys Cys Pro Xaa Leu Cys Leu Phe Phe Ser Phe Cys 1 5 10 15
20	The Ser Ser Gly Ser Cys Pro Phe Ser His Val Ser Gln Leu Ser Phe 20 25 30
30	Ile Ala Thr Phe Ser Gln Ser Ser Pro Val Leu Leu Val Pro Ala Tyr 35 40 45
35	Asn Thr Tyr Leu Ser Phe Leu Ala Phe Leu Asp Cys Ala Ser Leu Thr 50 55 60
	Ser Thr Xaa 65
40	
٠.	(2) INFORMATION FOR SEQ ID NO: 241:
45	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 69 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 241:
50	Met Ser Thr Phe Gln Leu Leu Leu Leu Ile Leu Ala Gln Ser Thr Tyr 1 5 10 15
55	Lys Ile Lys Ser Lys Pro Leu His Met Thr Asn His Thr Leu Leu Asn 20 25 30
55	Ser Pro Gly Leu Asn Pro Ser Ser Pro Thr Leu Asn Phe Lys Thr Gln 35 40 45
60	Gln His Glu Ser Val Ser Tyr Ala Cys Cys His Met Arg Ser Leu His 50 55 60

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His Ala Phe Ala Xaa
      65
 5
      (2) INFORMATION FOR SEQ ID NO: 242:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 44 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 242:
      Met Val Ser Val Val Leu Ile Phe Ser Phe Leu Ser Leu Thr Ile Ser
15
                                         10
      Thr Thr Ala Ser Ala Tyr Asn Gly Asn Asp Thr Gln Gly Trp Asn Asp
20
      Lys Phe His Kaa Kaa Ser Val Lys Thr Gln Thr Kaa
                                   40
               35
25
      (2) INFORMATION FOR SEQ ID NO: 243:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 51 amino acids
                     (B) TYPE: amino acid
30
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 243:
      Met Ile Ser Asp Ala Gly Ala Gly Phe Gly Val Phe Leu Leu Val Pro
 35
      Arg Ala Gly His Cys Trp Gly Ala Gly Lys Pro Leu Pro Ser Cys Pro
       Ser Val Ala Ser Ile Pro Ser Trp Val Leu Pro Ser Phe Leu Glu Arg
 40
                                   40
       Gly Arg Xaa
           50
 45
       (2) INFORMATION FOR SEQ ID NO: 244:
               (i) SEQUENCE CHARACTERISTICS:
 50
                      (A) LENGTH: 43 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 244:
 55
       Met Val Gln Thr Ile Gln Asp Phe Leu Ser Leu Phe Ser Thr Pro Ile
                                           10
       Phe Leu Leu Leu Met Phe Glu Thr Leu Ser Leu Ala Pro Ala Trp
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Leu Lys Pro Leu Arg Val Thr Ser His Ser Zas
                      . 40
              35
5
     (2) INFORMATION FOR SEQ ID NO: 245:
             (i) SEQUENCE CERFACIERISTICS:
                    (A) LEWIH: 61 amino acids
10
                    (B) TYFE: amino acid
                    (D) TOPILUSY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 245:
     Met Ile Leu Met Pro Gly Leu Gly Thr Ser Arg Gln Arg Ser Val Pro
15
      Phe Val Pro Thr Leu Asm Ala Ser Thr Pro Gly Ala Met Thr Gly Pro
                   20
20
      Thr Ala Thr Leu Thr Ser Dys Gln Trp Thr Thr Ala Cys Arg Val Ser
      Trp Ala Asn Gly Trp Tir Ser Let Arg Thr Phe Arg Xaa
25
                              55
      (2) INFORMATION FOR SEQ ID NO: 245:
30
              (i) SEQUENCE CERFACTERISTICS:
                     (A) LEWIH: 36 amino acids
                     (B) TYFE: amino acid
                     (D) TOPCLOGY: Linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 246:
35
      Met Ser His His Ala Glm Pro Arg Phe Leu Leu Ile Thr Met Leu Leu
                                           10
        1
       Gln Glu Ala Lys Pro Val Ser Asn Ile Pro His Leu Leu Glu Ser Trp
 40
       Tyr Phe Gly Xaa
               35
 45
      (2) INFORMATION FOR SEQ ID NO: 247:
               (i) SEQUENCE CFARACTERISTICS:
 50
                      (A) LEXTE: 33 amino acids
                      (B) T:::: amino acid
                      (D) TOPOLOGY: Linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 247:
 55
       Met Asn Ser Leu Phe Trp Met Ile Le: Leu Pro Val Ser Gln Asp Gln
```

Val Val Glu Gly Leu Glm Gly Rly Phe Ser Glm Ile His Met Arg Ile

Leu Arg Lys His Leu Xaa 35

5

(2)	INFORMATION	FOR	SEO	ID	NO:	248:
121	INFURMATION	FUN	2			

(i) SEQUENCE CHARACTERISTICS:

10

- (A) LENGTH: 211 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 248:
- Met Ser Arg Ser Xaa Asp Val Thr Asn Thr Thr Phe Leu Leu Met Ala 10

Ala Ser Ile Tyr Leu His Asp Gln Asn Pro Asp Ala Ala Leu Arg Ala 20

20

Leu His Gln Gly Asp Ser Leu Glu Cys Thr Ala Met Thr Val Gln Ile 40

- Leu Leu Lys Leu Asp Arg Leu Asp Leu Ala Arg Lys Glu Leu Lys Arg 25 55
 - Met Gln Asp Leu Asp Glu Asp Ala Thr Leu Thr Gln Leu Ala Thr Ala
- Trp Val Ser Leu Ala Thr Gly Gly Glu Lys Leu Gln Asp Ala Tyr Tyr 30 90
 - Ile Phe Gln Glu Met Ala Asp Lys Cys Ser Pro Thr Leu Leu Leu 105

35

- Asn Gly Gln Ala Ala Cys His Met Ala Gln Gly Arg Trp Glu Ala Ala
- Glu Gly Leu Leu Gln Glu Ala Leu Asp Lys Asp Ser Gly Tyr Pro Glu 40 135
 - Thr Leu Val Asn Leu Ile Val Leu Ser Gln His Leu Gly Lys Pro Pro 150
- Glu Val Thr Asn Arg Tyr Leu Ser Gln Leu Lys Asp Ala His Arg Ser 45 170
 - His Pro Phe Ile Lys Glu Tyr Gln Ala Lys Glu Asn Asp Phe Asp Arg 185 180

50

Leu Val Leu Gln Tyr Ala Pro Ser Ala Glu Ala Gly Pro Glu Leu Ser 205

Gly Pro Xaa 55 210

(2) INFORMATION FOR SEQ ID NO: 249:

			11, 0					18 am		acid	ls					
5			(xi)) TC	POLC	GY:	line	ar	QII	NO:	249	:			
	Met 1	Glu	Asp	Ser	Glu 5	Ala	Leu	Gly	Phe	Glu 10	His 1	Met (Gly :	Leu i	Asp 1	Pro
10	Arg	Leu	Leu	Gln 20	Ala	Val	Thr	Asp	Leu 25	Gly	Trp	Ser i	Arg	Pro' 30	Thr :	Leu
15	Ile	Gln	Glu 35	Lys	Ala	Ile	Pro	Leu 40		Leu	Glu	Gly I	Lys 45	Asp :	Leu	Leu
13	Ala	Arg 50		Arg	Thr	Gly	Ser 55	Gly	Lys	Thr	Ala	Ala 60	Tyr	Ala	Ile	Pro
20	Met 65	Leu	Gln	Leu	Leu	Leu 70	His	Arg	Lys	Ala	Thr 75	Gly	Pro	Val	Val	Glu 80
	Gln	Ala	Val	Arg	Gly 85	Leu	Val	Leu	Val	Pro 90	Thr	Lys	Glu	Leu	Ala 95	Arg
25	Gln	Ala	Gln	Ser 100	Met	Ile	Gln	Gln	Leu 105	Ala	Thr	Tyr	Cys	Ala 110	Arg	Asp
30			115					120					125			
	Ala	Val		Met	Glu	Lys	Pro 135		Val	Val	Val	Gly 140	Thr	Pro	Ser	Arg
35	145			His		150					155					160
	Glu	Let	ı Lev	ı Val	Val 165		Glu	, Ala	. Asp	Leu 170		Phe	Ser	Phe	Gly 175	Phe
40	Glu	Gl	ı Glu	1 Leu 180		Ser	Leu	ı Leu	185		: Leu	Pro	Arg	Ile 190		Gln
45	Ala	A Pho	e Le:		: Ser	Ala	. Thi	200		Glu	ı Asp	Val	Gln 205		Leu	Lys
	Glu	1 Le 21		e Lev	ı His	: Ası	219	_	Thr	Leu	ı Lys	220		Glu	. Ser	Gln
50	Let 225		o Gl	y Pro) Ası	230		u Glr	ı Glr	n Ph€	235		. Val	. Cys	: Glu	240
	Gl	u Gl	u As	p Ly:	24:		ı Le	u Lei	ı Tyı	250		ı Lev	Lys	Leu	255	Leu i
55	11	e Ar	g Gl	y Ly: 26		r Le	u Le	u Pho	e Va 26		n Thi	r Lev	ı Glu	270		Tyr
60	Ar	g Le	u Ar 27		u Ph	e Le	u Gl	u Gl: 28		e Se	r Ile	e Pro	28:		s Vai	l Leu

•	Asn Gly Glu Leu Pro Leu Arg Ser Arg Cys His Ile Ile Ser Gln Phe 290 295 300
5	Asn Gln Gly Phe Tyr Asp Cys Val Ile Ala Thr Asp Ala Glu Val Leu 305 310 315 320
	Gly Ala Pro Val Lys Gly Lys Arg Arg Gly Arg Gly Pro Lys Gly Asp 325 330 335
10	Lys Ala Ser Asp Pro Glu Ala Gly Val Ala Arg Gly Ile Asp Phe His 340 345 350
15	His Val Ser Ala Val Leu Asn Phe Asp Leu Pro Pro Thr Pro Glu Ala 355 360 365
13	Tyr Ile His Arg Ala Gly Arg Thr Ala Arg Ala Asn Asn Pro Gly Ile 370 . 375 380
20	Val Leu Thr Phe Val Leu Pro Thr Glu Gln Phe His Leu Gly Lys Ile 385 390 395 400
	Glu Glu Leu Leu Ser Gly Glu Asn Arg Gly Pro Ile Leu Leu Pro Tyr 405 410 415
25	Gln Phe Arg Met Glu Glu Ile Glu Gly Phe Arg Tyr Arg Cys Arg Asp 420 425 430
30	Ala Met Arg Ser Val Thr Lys Gln Ala Ile Arg Glu Ala Arg Leu Lys 435 440 445
	Glu Ile Lys Glu Glu Leu Leu His Ser Glu Lys Leu Lys Thr Tyr Phe 450 455 460 .
35	Glu Asp Asn Pro Arg Asp Leu Gln Leu Leu Arg His Asp Leu Pro Leu 465 470 475 480
	His Pro Ala Val Val Lys Pro His Leu Gly His Val Pro Asp Tyr Leu 485 490 495
40	Val Pro Pro Ala Leu Arg Gly Leu Val Arg Pro His Lys Lys Arg Lys 500 505 510
45	Lys Leu Ser Ser Ser Cys Arg Lys Ala Lys Arg Ala Lys Ser Gln Asn 515 520 525
	Pro Leu Arg Ser Phe Lys His Lys Gly Lys Lys Phe Arg Pro Thr Ala 530 535 540
50	Lys Pro Ser Xaa 545
55	(2) INFORMATION FOR SEQ ID NO: 250: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 299 amino acids
60	(B) TYPE: amino acid (D) TOPOLOGY: linear

	Met 1	Thr	Thr	Val	Pro 5	Pro	Ser	Pro	Arg	Pro 10	Met	Ser	Arg	Pro	Ser 15	Glu
5	Arg	Asn	Met	Arg 20	Arg	Pro	Arg	Gly	Pro 25	Ser	Pro	Leu	Pro	Ala 30	Ser	Pro
10	Arg	Asn	Ser 35	Thr	Pro	Asp	Glu	Pro 40	Asp	Val	His	Phe	Ser 45	Lys	ГЛЗ	Phe
ıo	Leu	Asn 50	Val	Phe	Met	Ser	Gly 55	Arg	Ser	Arg	Ser	Ser 60	Ser	Ala	Glu	Ser
15	Phe 65	Gly	Leu	Phe	Ser	Cys 70	Ile	Ile	Asn	Gly	Glu 75	Glu	Gln	Glu	Gln	Thr 80
	His	Arg	Ala	Ile	Phe 85	Arg	Phe	Val	Pro	Arg 90	His	Glu	Asp	Glu	Leu 95	Glu
20	Leu	Glu	Val	Asp 100	Asp	Pro	Leu	Leu	Val 105	Glu	Leu	Gln	Ala	Glu 110	Asp	Тут
25	Trp	Tyr	Glu 115		Tyr	Asn	Met	Arg 120	Thr	Gly	Ala	Arg	Gly 125	Val	Phe	Pro
	Ala	Tyr 130		Ala	Ile	Glu	Val 135		Lys	Glu	Pro	Glu 140	His	Met	Ala	Ala
30	Leu 145		Lys	Asn	Ser	Asp 150		Val	Asp	Gln	Phe 155	Arg	Val	Lys	Phe	Leu 160
	Gly	Ser	Val	Gln	Val 165		Тут	His	Lys	Gly 170		Asp	Val	Leu	Cys 175	Ala
35	Ala	Met	Gln	Lys 180		Ala	Thr	Thr	Arg 185		Leu	Thr	Val	His 190	Phe	Asn
40	Pro	Pro	Ser 195		Cys	Val	Leu	Glu 200		Ser	· Val	Arg	Gly 205		Lys	Ile
	Gly	Val 210	_	Ala	Asp	Asp	Ser 215		Glu	. Ala	Lys	Gly 220		Lys	Cys	Ser
45	His 225		Phe	e Gln	Leu	Lys 230		Ile	: Ser	Phe	235		Туг	His	Pro	Lys 240
	Asn	Asr	Lys	туг	Phe 245	-	Phe	lle	Thr	Lys 250		Pro	Ala	Asp	His 255	Arg
50	Phe	ala	ı Cys	His 260		. Phe	· Val	. Ser	Glu 265		Ser	Thr	Lys	270		Ala
55	Glu	ı Sei	275	-	/ Arg	, Ala	a Phe	Glr 280		Phe	е Тух	: Lys	Glr 285		· Val	Glu
-	Тут	Th:	_	s Pro	Thr	Glu	1 Asp 299		тут	Let	ı Glu	1				

	(2)	INFO	RMAT	NOI	FOR	SEQ	ID N	O: 2	51:							
5			(i) S (xi)	(1 (1 (1	A) LI B) T C) T(ENGTI YPE : OPOLA	H: 40 amin OGY:	o ami no ac line	ino a cid ear	acid		: 25 1	L:			
10	Leu 1	Leu	Tyr	Leu	Leu 5	Lys	Val	Xaa	Val	Ile 10	Phe	Val	Phe	Ser	Ser 15	Ser
	Lys	Gly	Val	Thr 20	Leu	Val	Ser	Met	Asn 25	Leu	Thr	Ser	Phe	Phe 30	Val	Ser
15	Ser	Val	Leu 35	Ala	Суз	Phe	Ser	Хаа 40								
20	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	7O: 2	:52 :							
			(i) S	_			RACTI H: 5				ds					
25			(xi)	()	B) T D) T	YPE: OPOL	amii OGY:	no a	cid ear			: 25	2:			
30	Met 1	Pro	Ala	Ser	Ser 5	Leu	Glu	Ser	Arg	Ser 10	Phe	Leu	Leu	Ala	Lys 15	Lys
50	Ser	Gly	Glu	Asn 20	Val	Ala	Lys	Phe	Ile 25	Ile	Asn	Ser	Tyr	Pro 30	Lys	Tyr
35	Phe	Gln	Lys 35	Asp	Ile	Ala	Glu	Pro 40	His	Ile	Pro	Cys	Leu 45	Met	Pro	Glu
	Tyr	Phe 50	Glu	Pro	Gln	Ile	Lys 55	Asp	Ile	Ser	Glu	Ala 60	Ala	Leu	Lys	Glu
40	Arg 65		Glu	Leu	Arg	Lys 70	Val	Lys	Ala	Ser	Val 75	Asp	Met	Phe	Asp	Gln 80
45	Leu	Leu	Gln	Ala	Gly 85	Thr	Thr	Val	Ser	Leu 90	Glu	Thr	Thr	Asn	Ser 95	Leu
	Leu	Asp	Xaa	Leu 100	Cys	Tyr	Tyr	Gly	Asp 105	Gln	Glu	Pro	Ser	Thr 110	Asp	Tyr
50	His	Phe	Gln 115	Gln	Thr	Gly	Gln	Ser 120	Glu	Ala	Leu	Glu	Glu 125	Glu	Asn	Asp
	Glu	Thr 130	Ser	Arg	Arg	Lys	Ala 135	Gly	His	Gln	Phe	Gly 140	Val	Thr	Trp	Arg
55	Ala 145	_	Asn	Asn	Ala	Glu 150	_	Ile	Phe	Ser	Leu 155	Met	Pro	Glu	Lys	Asn 160
	Glu	His	Ser	Tyr	Cys 165		Met	Ile	Arg	Gly 170	Met	Val	Lys	His	Arg 175	

	Tyr	Glu	Gln	Ala 180	Leu	Asn	Leu	Tyr	Thr 185	Glu	Leu	Leu	Asn	Asn 190	Arg	Leu
5	His	Ala	Asp 195	Val	Tyr	Thr	Phe	Asn 200	Ala	Leu	Ile		Ala 205	Thr	Val	Cys
	Ala	Ile 210	Asn	Glu	Lys	Phe	Glu 215	Glu	Lys	Trp	Ser	Lys 220	Ile	Leu	Glu	Leu
10	Leu 225	Arg	His	Met	Val	Ala 230	Gln	Lys	Val	Lys	Pro 235	Asn	Leu	Gln	Thr	Phe 240
15	Asn	Thr	Ile	Leu	Lys 245	Cys	Leu	Arg	Arg	Phe 250	His	Val	Phe	Ala	Arg 255	Ser
13	Pro	Ala	Leu	Gln 260	Val	Leu	Arg	Glu	Met 265	Lys	Ala	Ile	Gly	Ile 270	Glu	Pro
20	Ser	Leu	Ala 275	Thr	Туг	Ĥis	His	Ile 280	Ile	Arg	Leu	Phe	Asp 285		Pro	Gly
	Asp	290		Lys	Arg	Ser	Ser 295		Ile	Ile	Tyr	Asp 300	Ile	Met	Asn	Glu
25	Leu 305		Gly	. Lys	Arg	Phe 310		Pro	Lys	Ąsp	Pro 315	Asp	Asp	Asp	Lys	Phe 320
30	Phe	Glr	ı Ser	Ala	Met 325		Ile	: Cys	Ser	Ser 330		Arg	Asp	Leu	Glu 335	Leu
50	Ala	туз	c Glr	1 Val 340		Gly	Leu	. Leu	1 Lys 345		Gly	Asp	Asn	350		Phe
35	Ile	e Gly	7 Pro		Glr	His	Arg	Asn 360		туг	Туг	Ser	369		Phe	Asp
	Let	1 Ile 37		s Lev	ı Met	Glu	Glr 375		e Asp	Val	Thr	Leu 380	Lys	Trp	туг	Glu
40	As ₁		u Il	e Pro	Sei	390		r Phe	e Pro	His	395		Thi	r Met	: Ile	His 400
45	Le	ı Le	u Gli	n Ala	40!		Va.	l Ala	a Ası	1 Arg		Glu	ı Va	l Ile	415	Lys 5
73	11	e Tr	p Ly	s Ası 420		r Ly:	s Gl	ц Туг	42!	y Hi: 5	s Thi	Phe	a Ar	g Se:	r Ası) Leu
50	Ar	g Gl	u Gl 43		e Le	u Me	t Le	u Me 44		a Ar	g Ası) Lys	44		o Pro	o Glu
	Le	u Gl 45		l Al	a Ph	e Ala	a As 45		s Al	a Al	a Asp	9 Ile 460		s Se	r Al	а Туг
55	G1 46		er Gl	n Pr	o Il	e Ar 47		n Th	r Al	a Gl	n Ası 47		p Pr	o Al	a Th	r Ser 480
60	Le	u As	sn C)	/s Il	e Al 48		e Le	eu Ph	e Le	u Ar 49		a Gl	y Ar	g Th	r Gl 49	n Glu. 5

•	Ala	Trp	Lys	Met 500	Leu	Gly	Leu	Phe	Arg 505	Lys	His	Asn	Lys	Ile 510	Pro	Arg
5	Ser	Glu	Leu 515	Leu	Asn	Glu	Leu	Met 520	Asp	Ser	Ala	Lys	Val 525	Ser	Asn	Ser
	Pro	Ser 530	Gln	Ala	Ile	Glu	Val 535	Val	Glu	Leu	Ala	Ser 540	Ala	Phe	Ser	Leu
10	Pro 545	Ile	Cys	Glu	Gly	Leu 550	Thr	Gln	Arg	Val	Met 555	Ser	Asp	Phe	Ala	Ile 560
15	Asn	Gln	Glu	Gln	Lys 565	Glu	Ala	Leu	Ser	Asn 570	Leu	Thr	Ala	Leu	Thr 575	Ser
.5	Asp	Ser	Asp	Thr 580	Asp	Ser	Ser	Ser	Asp 585	Ser	Asp	Ser	Asp	Thr 590	Ser	Glu
20	Gly	Lys														
	(2)	INF	ORMA	TION	FOR	SEO	ID 1	NO: :	253:							
25	ν-,		•	SEQU	ENCE	СНА	RACT	ERIS	TICS		.ds					
				((B) 1	YPE:	ami	no a	cid							
				1	(D) 1	OPOL	OGY:	111	ıear							
30			(xi)		(D) 1 OKBU					EQ I	D NO	: 25	3:			
30	Met 1	_		SEC	UENC	E DE Cys	SCRI	PTIC	N:S		Ala			Pro	Leu 15	Leu
30 35	1		Leu	SEC Asn	Leu 5 Gln	E DE	SCRI Ile	Pro	N: S Asn	Trp 10	Ala	Arg	Cys		15 Asp	
35	1 Leu	. Leu	Leu Phe	SEQ Asn Pro 20	Leu 5 Gln	E DE Cys Leu	Ile Leu	PTIC Pro	Asn Phe 25	Trp 10	Ala Gly	Arg	Cys	Asp 30	15 Asp	
	1 Leu Leu	Leu Lys	Leu Phe Ala 35	SEQ Asn Pro 20	Leu 5 Gln	Cys Leu	Ile Leu Asn	PTIC Pro Pro Leu 40	Asn Phe 25	Trp 10 Glr	Ala Gly	Arg	Cys Asp Pro 45	Asp 30 Trp	15 Asp Gly	Pro
35	Leu Leu Lys	Leu Lys Ala 50	Phe Ala	SEQ Asn Pro 20 Lys	Leu 5 Gln : Ala	Cys Leu Ala	Ile Leu Asn Val	Pro Pro Leu 40	Asn Phe 25 Val	Trp 10 Glr Glu	Ala Gly Ala Val	Arg Glu Val Arg 60	Cys Asp Pro 45	Asp 30 Trp	Asp Gly	Pro Ile
35	Leu Lys Ser 65	Leu Lys Ala 50 Cys	Leu Phe Ala 35 A Pro	SEQ SEQ SEC	Leu 5 Gln Ala : Ala	Cys Leu Ala Glm	Leu Asm Val	Pro Pro Pro A0 Thr	Asn Phee 25 Val	Trp 10 Glr Glr Let	Ala Gly Ala Val Val Lev 75	Arg	Cys Asp Pro 45 Val	Asp 30 Trp Gln	Asp Gly Leu	Pro Ile Gln Ser 80
35	Leu Lys Ser 65	Lys Ala 50 Cys	Phe Phe Pro	SEC Asn Pro 20 20 Lys	Leu 5 Gln Ala Phe Ser 85	Cys Leu Ala Glm 70	Leu Asn Val 55	Pro Pro Pro Au	Asn Phe 25 Val Cys	Trp 10 Glr. Glr. Leu Leu 90	Ala Gly Ala Val Val 75	Arg	Pro 45	Asp 30 Trp	Asp Gly Gly Glr Glr Tyr	Pro Ile Gln Ser 80
35 40 45	Leu Lys Ser 65	Leu Lys Ala 50 Cys Gly	Phe Phe 35 Thr	SEC Asn Asn Pro 20 Lys Ser Pro Ser 116 100 Let Let	Leu 5 6 Gln 7 Phe 8 Ser 8 Ser 8 Ser	Cys Leu Ala Glm Cys	Leu Lau Val SS Pro	Pro Pro Pro Pro A0 Thr	Asn Phee 25 Val Cys Thr	Trp 10 Glr. Glr. Let	Ala Gly Ala Ala Val 1 Lev 75	Arg	Pro 45 Val	Asp 30 Trp Gln Ser Pro	Asp Gly Gly Glr	Pro Ile Gln Ser 80

•	(2) INFORMATION FOR SEQ ID NO: 254:	
5	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 21 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 254: 	
10	Met Arg Tyr His Ala Gln Leu Ile Phe Cys Ile Phe Cys Xaa Phe Val 1 5 10 15	
	Phe Val Xaa Lys Xaa 20	
15		
	(2) INFORMATION FOR SEQ ID NO: 255:	
20	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 255: 	
25	Met Asn Asp Asn Ser Pro Asn His Ser Ser Ser Tyr Leu Pro Leu Pro 1 5 10 15	
30	Leu Thr Ile Val Ile Leu Gln Thr Gly His Lys Gly Thr Leu Xaa 20 25 30	
	(2) INFORMATION FOR SEQ ID NO: 256:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 219 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 256:	
	Met His Phe Leu Phe Arg Phe Ile Val Phe Phe Tyr Leu Trp Gly Leu 1 5 10 15	
45	Phe Thr Ala Gln Arg Gln Lys Lys Glu Glu Ser Thr Glu Glu Val Lys 20 25 30	
	Ile Glu Val Leu His Arg Pro Glu Asn Cys Ser Lys Thr Ser Lys Lys 35 40 45	
50	Gly Asp Leu Leu Asn Ala His Tyr Asp Gly Tyr Leu Ala Lys Asp Gly 50 55 60	-
55	Ser Lys Phe Tyr Cys Ser Arg Thr Gln Asn Glu Gly His Pro Lys Trp 65 70 75 80	
	Phe Val Leu Gly Val Gly Gln Val Ile Lys Gly Leu Asp Ile Ala Met 85 90 95	
60	Thr Asp Met Cys Pro Gly Glu Lys Arg Lys Val Val Ile Pro Pro Ser	

	Phe	Ala	Tyr 115	Gly	Lys	Glu	Gly	Tyr 120	Ala	Glu	Gly	Lys	Ile 125	Pro	Pro	Asp
5	Ala	Thr 130	Leu	Ile	Phe	Glu	Ile 135	Glu	Leu	Tyr	Ala	Val 140	Thr	Lys	Gly	Pro
10	Arg 145	Ser	Ile	Glu	Thr	Phe 150	Lys	Gln	Ile	Asp	Met 155	Asp	Asn	Asp	Arg	Gln 160
	Leu	Ser	Lys	Ala	Glu 165	Ile	Asn	Leu	Туг	Leu 170	Gln	Arg	Glu	Phe	Glu 175	Lys
15	Asp	Glu	Lys	Pro 180	Arg	Asp	Lys	Ser	Туг 185	Gln	Asp	Ala	Val	Leu 190	Glu	Asp
	Ile	Phe	Lys 195	Lys	Asn	Asp	His	Asp 200	Gly	Asp	Gly	Phe	Ile 205	Ser	Pro	Lys
20	Glu	Tyr 210	Asn	Val	Tyr	Gln	His 215	Asp	Glu	Leu	Xaa					
25	(2)	INFO	ORMA	rion	FOR	SEQ	ID 1	NO: 2	257 :							
			(i) :	SEQUI												
30			(xi)	C	B) T D) T	ENGT YPE: OPOL E DE:	ami OGY:	no a	cid ear			: 25	7 :			
		Trp	Val	Ile	Arg	Val	Phe	Gln	Lys	Thr	Phe	Leu	Phe	Phe	Val	Leu
35 .	1				5					10					15	
	Phe	Trp	Ser	Val 20	His	Cys	Ile	Ser	Asp 25	Lys	Phe	Gly	Cys	Leu 30	Trp	His
10	Val	Cys	Met 35	Lys	Arg	Glu	Gly	Asp 40	Xaa	Asn	Cys	Leu	Ser 45	Phe	Ser	Xaa
•	Leu	Xaa 50														
1 5																
	(2)	INFO	ORMAI	CION	POR	SEQ	ID N	Ю: 2	58:							
50				(1 (1	A) Li B) T D) T	ENGT YPE: OPOLA	H: 1: amii CGY:	22 ar no ac line	mino cid ear	acio						
				SEQU												
55	Met 1	Pro	Ser	Gln	Thr 5	Glu	Xaa	Phe	Ala	Ala 10	Суз	Gly	Gly	His	Ser 15	Leu
60	Leu	Leu	Val	Xaa 20	Leu	Pro	Leu	Gly	Leu 25	Pro	Phe	Cys	Pro	Arg 30	Ala	Ala.

	Leu Cys Asp Leu Pro Phe Ser Leu Pro Ser Phe Pro Gly Gln Ala Arg 35 40 45
5	Arg Gly Gly Ala Glu Lys Gln Gly Ala Glu Gly Arg Gly Leu Gln Val 50 55 60
	Lys Pro Arg Gly Gln Arg Thr Phe Gln Val Ser Arg Thr Ala Pro Ala 65 70 75 80
10	Ala Pro Arg Ser Arg Gln Pro Arg Pro Pro Ala Ala Leu Pro Ala Leu 85 90 95
15	Gly Phe Gly Gly Arg Gly Val Ala Lys Gly Arg Phe Leu Cys Phe Trp 100 105 110
	Cys Leu Tyr Met Leu Arg Ile Asp Gln Xaa 115 120
20	(2) INFORMATION FOR SEQ ID NO: 259:
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 88 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 259:
30	Met Thr Ala Phe Cys Ser Leu Leu Leu Gln Ala Gln Ser Leu Leu Pro 1 5 . 10 15
	Arg Thr Met Ala Ala Pro Gln Asp Ser Leu Arg Pro Gly Glu Asp 20 . 25 30
35	Glu Gly Met Gln Leu Leu Gln Thr Lys Asp Ser Met Ala Lys Gly Ala 35 40 45
40	Arg Pro Gly Ala Xaa Arg Gly Arg Ala Arg Trp Gly Leu Ala Tyr Thr 50 55 60
	Leu Leu His Asn Pro Thr Leu Gln Val Phe Arg Lys Thr Ala Leu Leu 65 70 75 80
45	Gly Ala Asn Gly Ala Gln Pro Kaa 85
50	(2) INFORMATION FOR SEQ ID NO: 260:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 26 amino acids (B) TYPE: amino acid
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 260:
	Met Ile Gln Val Ser Val Pro Leu Leu Thr Ile Met Ile Phe Leu Leu 1 5 10 15
60	Tyr Leu Gln Ile Gly Pro Gly Lys Leu Xaa

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5 (2) INFORMATION FOR SEQ ID NO: 261: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 29 amino acids (B) TYPE: amino acid 10 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 261: Met Leu Leu Asp Pro Phe Ile Leu Leu Phe Cys Leu Phe Ser Thr Ala 10 15 Ala Gln Ser Cys Leu Glu Phe Ile Tyr Ile Gln Phe Xaa 20 20 (2) INFORMATION FOR SEQ ID NO: 262: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids 25 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 262: 30

Met Lys Phe Leu Ser Ile Leu Leu Asp Asp Asn Asn Phe Xaa Leu Met

Leu Met Leu Ala Pro Phe Gly Cys Leu Ala Phe Glu Arg Ser Met Lys 25

- Met Arg Asn Gly Ala Leu Gly Leu Glu Glu Val Xaa 35 40
- 40 (2) INFORMATION FOR SEQ ID NO: 263:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 363 amino acids
 - (B) TYPE: amino acid
- 45 (D) TOPOLOGY: linear

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 263:

Met Arg Thr Leu Phe Asn Leu Leu Trp Leu Ala Leu Ala Cys Ser Pro

Val His Thr Thr Leu Ser Lys Ser Asp Ala Lys Lys Ala Ala Ser Lys 25

Thr Leu Leu Glu Lys Ser Gln Phe Ser Asp Lys Pro Val Gln Asp Arg 55

Gly Leu Val Val Thr Asp Leu Lys Ala Glu Ser Val Val Leu Glu His 55 50

60 Arg Ser Tyr Cys Ser Ala Lys Ala Arg Asp Arg His Phe Ala Gly Asp

	65					70					75					80
5	Val	Leu	Gly	Tyr	Val 85	Thr	Pro	Trp	Asn	Ser 90	His	Gly	Tyr	Asp	Val 95	Thr
,	Lys	Val	Phe	Gly 100	Ser	Lys	Phe	Thr	Gln 105	Ile	Ser	Pro	Val	Trp 110	Leu	Gln
10	Leu	Lys	Arg 115	Arg	Gly	Arg	Glu	Met 120	Phe	Glu	Val	Thr	Gly 125	Leu	His	Asp
	Val	Asp 130	Gln	Gly	Trp	Met	Arg 135	Ala	Val	Arg	Lys	His 140	Ala	Lys	Gly	Leu
15	His 145	Ile	Val	Pro	Arg	Leu 150	Leu	Phe	Glu	Asp	Trp 155	Thr	Tyr	Asp	Asp	Phe 160
20	Arg	Asn	Val	Leu	Asp 165	Ser	Glu	Asp	Glu	Ile 170	Glu	Glu	Leu	Ser	Lys 175	Thr
20	Val	Va1	Gln	Val 180	Ala	Lys	Asn	Gln	His 185	Phe	Asp	Gly	Phe	Val 190	Val	Glu
25	Val	Trp	Asn 195	Gln	Leu	Leu	Ser	Gln 200	Lys	Arg	Val	Thr	Asp 205	Gln	Leu	Gly
	Met	Phe 210	Thr	His	Lys	Glu	Phe 215	Glu	Gln	Leu	Ala	Pro 220	Val	Leu	Asp	Gly
30	Phe 225	Ser	Leu	Met	Thr	Туг 230	Asp	Tyr	Ser	Thr	Ala 235	His	Gln	Pro	Gly	Pro 240
35	Asn	Ala	Pro	Leu	Ser 245	Trp	Val	Arg	Ala	Суs 250	Val	Gln	Val	Leu	Asp 255	Pro
•	Lys	Ser	Lys	Trp 260	_	Ser	Lys	Ile	Leu 265	Leu	Gly	Leu	Asn	Phe 270		Gly
40	Met	Asp	Tyr 275		Thr	Ser	Lys	Asp 280	Ala	Arg	Glu	Pro	Val 285	Val	Gly	Ala
	Arg	Tyr 290		Gln	Thr	Leu	Lys 295	_	His	Arg	Pro	Arg 300	Met	Val	Trp	Asp
45	Ser 305		Xaa	Ser	Glu	His 310	Phe	Phe	Glu	Tyr	Lys 315		Ser	Arg	Ser	Gly 320
50	Arg	His	Val	Val	Phe 325	-	Pro	Thr	Leu	Lys 330		Leu	Gln	Val	Arg 335	Leu
	Glu	Leu	Ala	Arg 340		Leu	Gly	Val	Gly 345		Ser	Ile	Trp	Glu 350		Gly
55	Gln	Gly	Leu 355	-	Tyr	Phe	Туг	Asp 360		Leu	Хаа					

			(i):	SEQU							٠.					
							H: l ami			acı	as					
							OGY:									
5			(xi)							EQ I	D NO	: 26	4:			
	Leu 1	Pro	Thr	Lys	Ile 5	Leu	Val	Lys	Pro	Asp 10	Arg	Thr	Phe	Glu	Ile 15	Lys
10	Ile	Gly	Gln	Pro 20	Thr	Val	Ser	Туг	Phe 25	Leu	Lys	Ala	Ala	Ala 30	Gly	Ile
15	Glu	Lys	Gly 35	Ala	Arg	Gln	Thr	Gly 40	Lys	Glu	Val	Ala	Gly 45	Leu	Val	Thr
	Leu	Lys 50	His	Val	Tyr	Glu	Ile 55	Ala	Arg	Ile	Lys	Ala 60	Gln	Asp	Glu	Ala
20	Phe 65	Ala	Leu	Gln	Asp	Val 70	Pro	Leu	Ser	Ser	Val 75	Val	Arg	Ser	Ile	Ile 80
	Gly	Ser	Ala	Arg	Ser 85	Leu	Gly	Ile	Arg	Val 90	Val	Lys	Asp	Leu	Ser 95	Ser
25	Glu	Glu	Leu	Ala 100	Ala	Phe	Gln	Lys	Glu 105	Arg	Ala	Ile	Phe	Leu 110	Ala	Ala
30	Gln	Lys	Glu 115	Ala	Asp	Leu	Ala	Ala 120	Gln	Glu	Glu	Ala	Ala 125	Lys	Lys	Xaa
35																
	(2)	INFO	ORMAI	SEQUI	ENCE	CHA	RACTI	ERIS	rics							
40			(xi)	()	B) T D) T	YPE: OPOL	H: 5 ami OGY: SCRI	no a lin	cid ear			: 26	5:			
45	Met 1	Leu	Leu	Gln	Ile 5	His	Pro	Leu	Leu	Pro 10	Ser	Pro	Thr	Ile	Pro 15	His
	Ile	Leu	Leu	Leu 20	Phe	Leu	Tyr	Pro	Thr 25	Phe	Ser	Ile	Leu	Glu 30	His	Ser
50	Cys	Ser	Tyr 35	Суз	Ile	Glu	Tyr	Leu 40	Trp	Val	Cys	Leu	Leu 45	Phe	Cys	Leu
55	Ser	Leu 50	Trp	Phe	Leu	Xaa										
-	(2)	INFO	ORMAI	MOI	FOR	SEQ	ID N	ю: 2	266:							
60			(i) :	SEQUI	ENCE	CHAI	RACTI	ERIS	rics	:						

```
(A) LENGTH: 29 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 266:
 5
      Met Cys Leu Trp Cys Cys Gly Asp Val Cys Ser Gly Leu Ser Ser Leu
      Leu Ser Leu Cys Val Cys Cys Val Val Leu Ala Val Cys
10
                   20
      (2) INFORMATION FOR SEQ ID NO: 267:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
20
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 267:
      Glu Gly Leu Arg Leu Leu Ser Leu Pro Ala Ala Leu Pro Arg Ser
                        5
25
      Cys Cys His Pro Arg Trp Leu Pro Val Xaa
                   20
30
      (2) INFORMATION FOR SEQ ID NO: 268:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 221 amino acids
                     (B) TYPE: amino acid
35
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 268:
      Met Phe His Gly Ile Pro Ala Thr Pro Gly Ile Gly Ala Pro Gly Asn
40
      Lys Pro Glu Leu Tyr Glu Glu Val Lys Leu Tyr Lys Asn Ala Arg Glu
                                      25
      Arg Glu Lys Tyr Asp Asn Met Ala Glu Leu Phe Ala Val Val Lys Thr
45
                                   40
      Met Gln Ala Leu Glu Lys Ala Tyr Ile Lys Asp Cys Val Ser Pro Ser
                               55
50
      Glu Tyr Thr Ala Ala Cys Ser Arg Leu Leu Val Gln Tyr Lys Ala Ala
                           70
      Phe Arg Gln Val Gln Gly Ser Glu Ile Ser Ser Ile Asp Glu Phe Cys
55
      Arg Lys Phe Arg Leu Asp Cys Pro Leu Ala Met Glu Arg Ile Lys Glu
                                      105
      Asp Arg Pro Ile Thr Ile Lys Asp Asp Lys Gly Asn Leu Asn Arg Cys
60
                                 120
```

	Ile Ala Asp Val Val Ser Leu Phe Ile Thr Val Met Asp Lys Leu Arg 130 135 140
5	Leu Glu Ile Arg Ala Met Asp Glu Ile Gln Pro Asp Leu Arg Glu Leu 145 150 155 160
10	Met Glu Thr Met His Arg Met Ser His Leu Pro Pro Asp Phe Glu Gly 165 170 175
10	Arg Gln Thr Val Ser Gln Trp Leu Gln Thr Leu Ser Gly Met Ser Ala 180 185 190
15	Ser Asp Glu Leu Asp Asp Ser Gln Val Arg Gln Met Leu Phe Asp Leu 195 200 205
	Glu Ser Ala Tyr Asn Ala Phe Asn Arg Phe Leu His Ala 210 215 220
20	·
	(2) INFORMATION FOR SEQ ID NO: 269:
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 3 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 269:
30	Met Lys Xaa 1
35	(2) INFORMATION FOR SEQ ID NO: 270:
40	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 49 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 270:
45	Met Gln Ala Pro Phe Xaa His Phe Ser Phe Arg Met Phe Ser Asn Lev 1 5 10 15
73	Tyr Cys Phe Ser Asp Phe Gln Pro Asn Ile Ser Pro Cys Pro Leu Cys 20 25 30
50	His Cys Ile Leu Pro Xaa His His His Val Phe Leu Leu Leu Ala Val 35 40 45
	Хаа
55	
	(2) INFORMATION FOR SEQ ID NO: 271:
60	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 52 amino acids

60

(2) INFORMATION FOR SEQ ID NO: 274:

	(B) TYPE: amino acid
	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 271:
5	Met Lys Leu Val Thr Met Phe Asp Lys Leu Ser Arg Asn Arg Val Ile 1 5 10 15
10	Gln Pro Met Gly Met Ser Pro Arg Gly His Leu Thr Ser Leu Gln Asp 20 . 25 30
10	Ala Met Cys Glu Thr Met Glu Gln Gln Leu Ser Ser Asp Pro Asp Ser 35 40 45
15	Asp Pro Asp Xaa 50
20	(2) INFORMATION FOR SEQ ID NO: 272: (i) SEQUENCE CHARACTERISTICS:
25	(A) LENGTH: 32 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 272:
	Met Ala Val Gly Glu Ala Val Phe Val Pro Leu Gln His Pro Pro Leu 1 5 10 15
30	Leu His Gly Ser Pro Ile Pro Lys Leu Leu Pro Gly Pro Leu Leu Xac 20 25 30
35	
40	(2) INFORMATION FOR SEQ ID NO: 273:
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 57 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 273:
45	Met Asn Gly Cys His Arg Arg Lys Arg Leu His Leu Cys Lys Thr Il. 1 5 10 15
50	Tyr Leu Leu Trp Phe Val Phe Ser Phe Leu Leu Ser Asn Glu Val Va 20 25 30
	Ser Ser His Trp His Ile Leu Arg Ala Val Gln Ile Ile Cys Thr Le 35 40 45
55	Phe His Arg Xaa Ile Ser Ala Phe Xaa 50 55

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(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 22 amino acids
                    (B) TYPE: amino acid
5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 274:
     Met Gly Trp Val Ser Ser Pro His Val Lys Arg Arg Glu Cys Val Leu
                                         10
                       5
10
     Lys Lys Pro Phe Phe Xaa
                  20
15
      (2) INFORMATION FOR SEQ ID NO: 275:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 275:
      Met Phe Asn Phe Phe Lys Asn Pro Leu Leu Thr Cys Leu Phe Ile Ser
25
                                          10
      Cys Tyr Leu Tyr Leu Ser Leu Leu Val Asn Lys Val Leu Phe Ala Glu
                   20
      Glu Gly Leu Cys Cys Thr Tyr Cys Thr Thr Ser Asn Thr Gly Glu Gly
30
                    40
              35
      Gly Val Xaa
           50
35
      (2) INFORMATION FOR SEQ ID NO: 276:
40
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 2 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 276:
45
      Met Xaa
        1
50
       (2) INFORMATION FOR SEQ ID NO: 277:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 66 amino acids
55
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 277:
      Met Leu Cys Thr Ile Leu Thr Val Val Ile Ile Ile Ala Ala Gln Thr
 60
                        5
                                           10
```

	Thr	Arg	Thr	Thr 20	Gly	Ile	Pro	Lys	Asn 25	Ala	Pro	Gly	Pro	Ala 30	Pro	Leu
5	Суз	Ala	Pro 35	Arg	Ser	Pro	Arg	Leu 40	Phe	Leu	Gln	Xaa	Tyr 45	Arg	Gly	Pro
10	Asn	Gly 50	Arg	Pro	Ala	His	Pro 55	Phe	Leu	Gly	Pro	Ser 60	Asp	Leu	Asp	Thr
	Ser 65	Xaa														
15	(2)	INF	ORMAT	rion	FOR	SEQ	ID 1	10: 2	278:							
20			(i) :	(ENGT	H: 2	57 a	TICS: mino cid		ds					
			(xi)	(D) TY	OPOL	OGY:	lin		EQ II	O NO	: 27	8:			
25	Met 1		Glý											His	Ser 15	Pro
	Gly	Leu	Pro	Leu 20	Val	Leu	Val	Leu	Leu 25	Ala	Leu	Gly	Ala	Gly	Trp	Ala
30	Gln	Glu	. Gly 35		Glu	Pro	Val	Leu 40	Leu	Glu	Gly	Glu	Cys 45	Leu	Val	Val
35	Cys	Glu 50	Pro	Gly	Arg	Ala	Ala 55		Gly	Gly	Pro	Gly 60	Gly	Ala	Ala	Leu
	Gly 65		Ala	Pro	Pro	Gly 70	Arg -		Ala	Phe	Хаа 75	Ala	Val	Arg	Ser	His 80
40			Glu		85					90					95	
	Тут	Phe	Asp	Gln 100		Leu	Val	Asn	Glu 105	Gly	Gly	Gly	Phe	Asp 110	Arg	Ala
45	Ser	Gly	/ Ser 115		Val	Ala	Pro	Val 120		Gly	Val	Туг	Ser 125		Arg	Phe
50	His	Va]		. Lys	Val	Tyr	Asn 135		Gln	Thr	Val	Gln 140		Ser	Leu	Met
	Let 145		1 Thr	Trp	Pro	Val 150		e Ser	Ala	Phe	Ala 155		Asp	Pro	Asp	Val 160
55	Thi	r Arg	g Glu	a Ala	165		Ser	Ser	Val	Leu 170		Pro	Leu	Asp	175	Gly
	Ası) Ar	y Val	180)	_			185	,				190	•	Leu
ራ በ	C1.		- T		. Dha	ton	. ~~		Dro	wie	In	Dre	. CA.	To) Acn

		195					200					205			
.	Pro Ser 210	Leu	Ser	Ser	Thr	Arg 215	Ile.	Gln	Pro	Leu	Thr 220	Thr	Phe	Phe	Cys
5	Pro Leu 225	Leu	⊋ro	Хаа	Lys 231	Gln	Хаа	Lys	Gln	Xaa 235	Хаа	Xaa	Ser	Leu	Trp 240
10	Leu Leu	Ser	His	Leu 245	Phe	Ala	طتق	Glu	Pro 250	Val	Pro	Asn	Thr	Gln 255	Val
	Xaa														
15															
	(2) ⊃ N F	OFMAC	MOI	FCP.	SEÇ	נ כנ	NC: 2	279:							
20		(i) :	(A) 1 B) T	eng: Ype:	H: 1 ami	EPIS 03 a no a lin	mino cid		ds					
		(xi)	SEÇ	UENC	E CE	SCRI	P TIC	N: 5	EQ I	D NO	: 27	9:			
25	Met Ala 1	220	Arg	Ala 5	Leu	520	GŢĀ	Ser	Ala 10	Val	Leu	Ala	Ala	Ala 15	Val
30	Phe Wal	Gly	Gly 20	λla	Val	Ser	Ser	Pro 25	Leu	Val	Ala	Pro	Asp 30	Asn	Gly
	Ser Ser	35	Thr	Leu	His	Ser	13 52	Thr	Glu	Thr	Thr	Pro 45	Ser	Pro	Ser
35	Asn Asr 50		Gly	Asn	Gly	His 55		Glu	Тут	Ile	Ala 60		Ala	Leu	Val
·	Pro Val	l Phe	Phe	Ile	Met 70		· Leu	Phe	Gly	Val 75		Ile	Xaa	Pro	Xaa 80
40	Хаа Заа	a Lys	Lys	Lys 85		Tyz	' ಸ್ವಾ	Cys	Thr 90		Glu	Ala	Glu	Gln 95	
45	Ilə Slu	ı Slu	Glu 100	-	Gly	Xaa	ı								
	(2) <u>IN</u>	FCRMA	TICH	FCR	seq	ID	NC:	280:							
50		(i)	_	(A)	leng:	r 4: :	TERIS 33 au iro a	nino		ds					
55		(xi)		(D)	ropci	COGY	: li	near	SEQ :	ED N	o: 21	30:			
,,	Met Pro	o 7al	Thr	Leu S		Ser	r Leu	Gly	Phe 10		Val	l Leu	ı Lev	Ser 19	
60	Leu Ph	e Pro	Trp 20		Th:	: Ası	slr	Gly 25		Gly	Pro	Ala	Thr 30		Tyr

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512

Xaa 5 (2) INFORMATION FOR SEQ ID NO: 281: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 281: 15 Met Val Leu Gly Leu Leu Leu Leu Leu Xaa Phe Phe Ser Phe Ser Ser 10 Ser Pro Ser Pro Ser Ser Ser Leu Leu Leu Ser Ser Phe Phe Phe 20 25 20 Gln Ser Leu Ala Leu Ser Pro Arg Leu Glu Xaa 35 40 25 (2) INFORMATION FOR SEQ ID NO: 282: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 21 amino acids 30 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 282: Glu Trp Leu Val Phe Thr Phe Leu Leu Val Phe Gly Ser Pro Leu Gly 35 5 10 Lys Gly Pro Leu Xaa 20 40 (2) INFORMATION FOR SEQ ID NO: 283: (i) SEQUENCE CHARACTERISTICS: 45 (A) LENGTH: 70 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 283: 50 Met Ile Arg Ala Leu Ser Leu Phe Leu Leu Ile Phe Asp Ala Ala Leu Phe Ser Leu Ser Val Phe Val Phe Ile Gly His Leu Leu Pro Met Pro 25 55 Lys Gly Thr Gly Leu His Ser Cys Ala Lys His Leu Ile Lys Ser Leu

40

55

60

Lys Glu Asn Val Leu Pro Leu Met Asn Tyr Pro Asp Cys Lys Leu Lys

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Ile Asn Ile Ser Pro Xaa
5
     (2) INFORMATION FOR SEQ ID NO: 284:
            (i) SEQUENCE CHARACTERISTICS:
10
                   (A) LENGTH: 75 amino acids
                   (B) TYPE: amino acid
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 284:
     Met Gly Lys Leu Ile Arg Leu Ser Val Met Val Met Ser Val Arg Arg
15
     Leu Phe Ser Ile Tyr Trp Val Leu Ser Thr Val Pro Asp Ala Val Gly
                                    25
                  20
20
     Ser Arg Gly Gly Met Glu Glu Cys Ser Arg Gly Leu Cys Cys Val
                                  40
     Ala Gly Gln His Lys Gln Ala Lys Gly Lys Arg Gln Ala Trp Asn Lys
25
                              55
      Gly Gly Glu Tyr Gln Cys Val Thr Tyr Cys Xaa
                          70
30
      (2) INFORMATION FOR SEQ ID NO: 285:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 33 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 285:
      Met Pro Ala Leu Val Thr Leu Leu Leu Phe Pro Leu Leu Pro Leu
40
              5
        1
      Met Glu Ala Ser Cys His Val Met Arg Cys Pro Met Glu Arg Pro Thr
                                25
                  20
45
      Xaa
50
      (2) INFORMATION FOR SEQ ID NO: 286:
              (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 17 amino acids
 55
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 286:
      Glu Ala Pro Trp Gly Leu Leu Lys Leu Leu Leu Leu Leu Ala Val Phe
 60
                                          10
           5
```

Xaa 5 (2) INFORMATION FOR SEQ ID NO: 287: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 17 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 287: 15 Met Gln Gln Lys Gln Lys Lys Ala Asn Glu Lys Lys Glu Glu Pro Lys 5 10 Xaa 20 (2) INFORMATION FOR SEQ ID NO: 288: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 38 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 288: 30 Met Gln Arg Lys Val Ser Asp Phe Ile Ile His Gln Arg Leu Thr Val Asn Leu Cys Val Ile Ser Phe Phe Phe Phe Leu Pro Ile Cys Ile Phe 35 20 25 Ser Leu Ala Lys Lys Xaa 35 40 (2) INFORMATION FOR SEQ ID NO: 289: (i) SEQUENCE CHARACTERISTICS: 45 (A) LENGTH: 12 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 289: 50 Met Ala Leu Leu Ile Ser Ser Leu Ile Trp Ser Xaa 1 5 10 55 (2) INFORMATION FOR SEQ ID NO: 290: (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 35 amino acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 290: Met Gln Met Phe Thr Val Ser Leu Leu Leu Ser Leu Leu Leu Arg Ser 5 Thr Asp Gln Asn His Leu Gln Leu Leu Val Gly Arg Glu Asp His Tyr 25 Gly Gly Xaa 10 (2) INFORMATION FOR SEQ ID NO: 291: 15 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 15 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 291: Met Ser Glu Ser Ala Cys Ile Leu Asn Asn Gln Lys Glu Leu Xaa 10 5 25 (2) INFORMATION FOR SEQ ID NO: 292: (i) SEQUENCE CHARACTERISTICS: 30 (A) LENGTH: 44 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 292: Met Asp Leu Asp Arg Val Lys Ala Glu Ala Thr Glu Asp Ile Thr Ser 35 Gly Val Leu Cys Leu Leu Phe Leu Arg Leu Pro Pro Asn Ser Cys Ile 20 25 40 Phe Pro Ser Ala Val Leu Gly Ser Thr Arg Thr Xaa 35 45 (2) INFORMATION FOR SEQ ID NO: 293: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 136 amino acids 50 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 293: Val Val Gly Thr Gly Thr Ser Leu Ala Leu Ser Ser Leu Leu Ser Leu 55 10 Leu Leu Phe Ala Gly Met Gln Met Tyr Ser Arg Gln Leu Ala Ser Thr 25 60 Glu Trp Leu Thr Ile Gln Gly Gly Leu Leu Gly Ser Gly Leu Phe Val WO 98/54963

	35	40	45	
5	Phe Ser Leu Thr Ala 50	Phe Asn Asn Leu Glu 55	Asn Leu Val 60	Phe Gly Lys
J	Gly Phe Gln Ala Lys 65	Ile Phe Pro Glu Ile 70	Leu Leu Cys 75	Leu Leu Leu 80
10	Ala Leu Phe Ala Ser	Gly Leu Ile His Arg		Thr Thr Cys 95
	Phe Ile Phe Ser Met 100	Val Gly Leu Tyr Tyr 105	lle Asn Lys	Ile Ser Ser 110
15	Thr Leu Tyr Gln Ala 115	Ala Ala Pro Val Leu 120	Thr Pro Ala 125	Lys Val Thr
20	Gly Lys Ser Lys Lys 130	Arg Asn Xaa 135		
	(2) INFORMATION FOR	SEQ ID NO: 294:		
25	(A) L (B) T	CHARACTERISTICS: ENGTH: 34 amino acid YPE: amino acid OPOLOGY: linear	ds	
30	(xi) SEQUENCI	E DESCRIPTION: SEQ	ID NO: 294:	
	Met Phe Ile Phe Leu 1 5	Phe Leu Cys Val Leu 10		Ile Gln Glu 15
35	Glu Tyr Tyr Arg Leu 20	Phe Lys Asn Val Pro 25	Cys Cys Phe	Gly Cys Leu 30
	Arg Xaa			
40				
	(2) INFORMATION FOR	SEQ ID NO: 295:		
45	(A) L (B) T (D) T	CHARACTERISTICS: ENGTH: 137 amino ac TYPE: amino acid OPOLOGY: linear E DESCRIPTION: SEQ		
50	Met Arg Thr Pro Gly	Pro Leu Pro Val Leo		Leu Ala Gly 15
55	Ala Pro Ala Ala Arg 20	Pro Thr Pro Pro Th	r Cys Tyr Ser	Arg Met Arg
<i></i>	Ala Leu Ser Gln Glu 35	Ile Thr Arg Asp Pho	e Asn Leu Leu 45	
60	Glu Pro Ser Glu Pro	Cys Val Arg Tyr Le	u Pro Arg Leu	Tyr Leu Asp

	Ile 65	His	Asn	Tyr	Суз	Val 70	Leu	Asp	Lys	Leu	Arg 75	Asp	Phe	Val	Ala	Ser 80
5	Pro	Pro	Cys	Trp	Lys 85	Val	Ala	Gln	Val	Asp 90	Ser	Leu	Lys	Asp	Lys 95	Ala
10	Arg	Lys	Leu	Tyr 100	Thr	Ile	Met	Asn	Ser 105	Phe	Cys	Arg	Arg	Asp 110	Leu	Val
	Phe	Leu	Leu 115	Asp	Asp	Cys	Asn	Ala 120	Leu	Glu	Tyr	Pro	Ile 125	Pro	Val	Thr
15	Thr	Val 130	Leu	Pro	Asp	Arg	Gln 135	Arg	Xaa							
20	(2)	INF		SEQU	ENCE	СНА	RACT	ERIS	TICS							
				(A) L B) T	YPE:	ami	no a	cid	acıd	S					
25			(xi)	SEQ	D) T					EQ I	D NO	: 29	6:			
	Met 1		Leu	Leu	Lys 5	Pro	Ser	Ala	His	Ser 10	Pro	Val	His	Xaa	Leu 15	Val
30	Leu	Leu	Phe	Pro 20	Arg	Gly	Trp	Ser	Gln 25	Pro	Gly	Thr	His	Lys 30	Arg	Gln
35	Ile	Leu	Val 35	Asn	Xaa	Ala	Ser	Leu 40	Pro	Gly	Gly	Cys	Leu 45	Leu	Pro	Trp
	Ile	Trp 50		Gly	Ala	Ala	Leu 55	Arg	Phe	Xaa						
40	(2)	INF	ORMA	tion	FOR	SEQ	ID	NO:	297 :							
			(i)	SEQU							1-					
45			(xi)	(A) L B) T D) T UENC	YPE:	ami OGY:	no a lin	cid ear			: 29	7:			
50	Met 1		Arg	Arg	Ala 5	Glu	Ala	Ser	Ile	Phe 10	Val	Leu	Pro	Lys	Thr 15	Leu
	Leu	Phe	Val	Leu 20		Pro	Ala	Phe	Pro 25		Pro	Ala	Val	Gly 30	Cys	Pro
55	Val	. Pro	Хаа 35													
60	(2)	TNE	אשברץ	ጥፐ∕ነኦ፣	פרק	gen.	ית	MO.	200.							

5			(i) ! (xi)	() () ()	A) L B) T D) T	ENGT YPE: OPOL	H: 7: ami: OGY:	8 am no a lin	ino d cid ear	acid		: 298	3:			
10	1	-	Tyr Gln	Phe	5				Val	10				Pro	15	
15	Ser	Val	Trp 35	20 Tyr	Glu	Arg	Tyr	Lys 40	25 Phe	Asp	Ile	Pro	Val 45	30 Phe	His	Leu
20		50	Gln				55					60		_	Leu	Glu
20	Lys 65	Gln	Leu	Leu	Lys	70	Glu	Gln	Gln	Ser	Thr 75	Gly	Xaa	Xaa		
25	(2)	INF	ORMA:			_										
30				(A) L B) T D) T	CHAI ENGT YPE: OPOL E DE	H: 9 ami OGY:	5 am no a lin	ino cid ear	acid		: 29	9:			
35	Met 1	Phe	Val	Leu	Phe 5	Ser	Leu	Pro	Lys	Tyr 10	Ala	Gly	Leu	Arg	Leu 15	Pro
	Ile	Pro	Gly	Leu 20	Ser	Ala	Leu	Leu	Val 25	Phe	Leu	Leu	Ser	Leu 30	Phe	Ser
40	Arg	Arg	Ala 35	Gln	Val	Glu	Leu	Thr 40	Thr	Gly	Arg	Glu	Thr 45	Leu	Pro	Lys
45	Asn	Leu 50	Gln	Gly	Tyr	Phe	Pro 55		Phe	Gly	Phe	Gln 60		Gln	Asn	Phe
45	Leu 65	Ser	Cys	Lys	Ile	Tyr 70	Ala	Ala	Ser	Gln	Lys 75	Gln	Pro	Leu	Pro	Pro 80
50	Leu	Tyr	· Gln	Leu	Arg 85		Tyr	Leu	Lys	His 90	Met	Gly	Leu	Pro	Хаа 95	
55	(2)	INF	ORMA			_										
				((A) I (B) T (D) T	CHA ENGI TYPE:	H: 4 ami OGY:	ino a	nino ncid near	ació						
60			(X7)	SEQ	ULINC	E DE	SCRI	PTIO	in:S	FQ I	אט	: 30	υ:			

	Met 1	Ser	Ser	His	Trp 5	Thr	Leu	Lys	Ile	Leu 10	Leu	Val	Pro	Leu	Phe 15	Tyr
5	Leu	Ser	Leu	Glu 20	Phe	Pro	Ser	GJY	Phe 25	Val	Leu	Cys	Leu	Ala 30	Asn	Asp
	Leu	Gly	Tyr . 35	His	Phe	Ser	Ser	Arg 40	Val	Arg	Ser	Xaa				
10																
	(2)	INF	ORMA!	rion	FOR	SEQ	ID 1	ю: 3	01:							
15				(A) L B) T D) T	engt YPE: OPOL	H: 3 ami OGY:	ERIST 1 am no ac line PTION	ino a cid ear	acid		. 30	1:			
20	Met							Leu						Phe	Ile	Phe
	1				5					10					15	
25	Leu	Суs	Tyr	Leu 20	Asp	Ala	Cys	Ile	Asn 25	Val	Phe	Суѕ	Phe	Tyr 30	Xaa	
	(2)	TATES	ODMA	ጥፐርእነ	EOB	SEO.	TD I	NO: 3	302.							
30	(2)	LINE				-		ERIS		:						
				((A) L	ENGI	H: 1	.13 a .no a	mino		.ds			-•		
25			(xi)					lin PTIO		EQ I	D NO	: 30	2:			
35	Met 1	Pro	Val	Leu	Pro	Gly	Arg	Thr			.	*	502			
	_		741		5			1111	Thr	Ala 10	ren	Leu	Ser	Leu	Thr 15	Leu
40		Phe			Pro		Ser	Gly		10 Glu					15	
40	Ala		Ala	Val 20	Pro	Cys			Val 25 Glu	10 Glu	Ala	Gly	Pro	Cys 30	15 Val	Pro
40 45	Ala Arg	Ser	Ala His	Val 20 Gly	Pro Cys	Cys	Ser	Gly Trp 40	Val 25 Glu	10 Glu Ala	Ala Ser	Gly Val	Pro Cys 45	Cys 30 Val	15 Val Thr	Pro Ser
45	Ala Arg Ser	Ser Thr 50	Ala His 35	Val 20 Gly	Pro Cys	Cys Ser Ser	Ser Trp 55	Gly Trp 40	Val 25 Glu Ala	Glu Ala Arg	Ala Ser	Gly Val Leu 60	Pro Cys 45	Cys 30 Val	Val Thr Ser	Pro Ser Ala
	Ala Arg Ser Ala	Ser Thr 50	His 35	Val 20 Gly Gly Arg	Pro Cys Gly	Ser Ser Ala	Ser Trp 55	Gly Trp 40	Val 25 Glu Ala Asp	Glu Ala Arg	Ala Ser Ala Pro 75	Gly Val Leu 60	Pro Cys 45 Phe	Cys 30 Val Pro	Val Thr Ser	Pro Ser Ala Gly 80 Gly
45	Ala Arg Ser Ala 65	Thr 50 Trr	His 35	Val 20 Gly Gly Arg	Pro Cys Gly Gly Gly S5 Ser	Cys Ser Ser Ala 70	Trp 55 Ala	Gly Trp 40 Arg	Val 25 Glu Ala Asp	Ala Arg	Ala Ser Ala Pro 75	Gly Val Leu 60 Trp	Pro Cys 45 Phe	Cys 30 Val Pro	Thr Ser Thr Gly 95	Pro Ser Ala Gly 80

```
(2) INFORMATION FOR SEQ ID NO: 303:
             (i) SEQUENCE CHARACTERISTICS:
 5
                    (A) LENGTH: 14 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 303:
10
      Thr His Ile His Thr His Ile Ile Cys Ser Ser Val Xaa
       1
                       5
                                          10
15
      (2) INFORMATION FOR SEQ ID NO: 304:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 35 amino acids
                    (B) TYPE: amino acid
20
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 304:
      Met Glu Asn Phe Phe Phe Ser Phe Tyr Leu Phe Leu Ile Thr Leu Ile
                                          10
25
      Pro Asn Gly Arg Thr Leu Ser Thr Thr Ala Asp His Cys Lys Ile Pro
                                      25
                   20
      Cys Ile Xaa
30
              35
      (2) INFORMATION FOR SEQ ID NO: 305:
35
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 35 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
40
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 305:
      Met Glu Leu Trp Glu Leu Ala Leu Cys Leu Leu Val Ala Leu Ser Ala
        1
                                           10
45
      His Met Phe Thr Val Gln Leu Leu Ala Asp Leu Gly Phe Leu Phe Gly
                                       25
      Gly Phe Xaa
               35
50
      (2) INFORMATION FOR SEQ ID NO: 306:
55
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 82 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 306:
60
```

	Met 1	Gly	Ala	Slγ	Ile E	Leu	λia	Leu	Leu	Leu 10	Pro	Leu	Glu	Ser	Val 15	Leu
5	Thr	Cys	Ser	729 26	Ile	Ser	Val	Ser	Thr 25	Ser	Glu	Arg	Gln	Leu 30	Trp	Gln
	Ser	Ser	Gln 35	Lys	Ala	The	Ile	Leu 40	Ser	Leu	Lys	Leu	Asp 45	Ser	Cys	Phe
10	Cys	Gly 50	His	Ser	Gly	Leu	Lys 55	Gly	Lys	Asn	Glu	Asp 60	Thr	Asp	Ser	Ser
15	Val 65	Pro	Ile	Ile	Pro	Ser 7)	Lys	Thr	His	Thr	His 75	Leu	Gly	Lys	His	Leu 80
	Ile	Xaa														
20	(2)	INFO	CEMPA	CION	FCR	SEQ	D:	xo: 3	307 :							
25				C	A) I 3) I	engt 172 : 070l	H: 7 ami OGY:	2 am no a lin	ino cid ear	acid		: 30	7:			
30	Met 1	Phe		Phe	Val 5	Leu	Phe	Ile	T/=	Ser 10	Ser	Ser	Glu	Thr	Trp 15	Ser
	Gĵλ	Ser	7al	Ala 20	Gln	ಸಿಕ್ತಾ	GŗÅ	Val	His 25	Gly	Val	Ile	Ile	Gly 30	His	Cys
35	Ser	Val	Glu 35	Leu	Pro	GŢĀ	Ser	Gly 40	ązĄ	Pro	Pro	Ala	Ser 45	Ala	Xaa	Leu
1 0	Val	Ala 50	Gly	Thr	Ile	Gly	Tile 55	Cys	Pro	Thr	Met	Pro 60	Gly	Phe	Val	Tyr
	Phe 65	Leu	Asn	λsp	Val	Ха <u>а</u> 70	Asn	Хаа								
1 5	(2)	INFO	ORMA:	CION	FCP.	SEQ	D I	NO: 3	308:							
50				C	A) L B) T D) T	enct YPE: OPOL	H: 3 ami CGY:	4 am no a lin	ino cid ear	acid		: 30	8:			
55	Met 1	Ąsp	Ser	The	Leu 5	A≃g	GĽn	Gly	Arg	Хаа 10	Leu	Leu	Thr	Leu	Val 15	Pro
	Ala	Ser	Leu	?he 20	Ser	Leu	Thr	Leu	Gly 25	Gly	Pro	Gly	Pro	Trp 30	Lys	Asp
50	Pro	Xaa														

,	(2)	INF	Jruna	LION	FUR	SEQ	נו עד	W: 3	.05:							
10				(A) Li B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	15 as no a lin	mino cid ear	aci		: 309) :			
15	Met 1	Gln	Val	Val	Gly 5	Ser	Trp	Pro	Gly	Arg 10	Val	Gly	Val	Val	Gly 15	Leu
-	Ala	Phe	Ser	Leu 20	Val	Ile	Pro	Pro	Pro 25	Ala	Ile	Cys	Ile	Ala 30	Gly	Pro
20	Ala	Pro	Gly 35	Leu	Gly	Gly	Gly	Glu 40	Arg	Gln	Gln	Lys	Gly 45	Leu	Gly	Arg
	Gly	Gly 50	Gly	Gly	Leu	Arg	Asn 55	Cys	Pro	Gly	Arg	Val 60	Gly	Met	Ala	Ala
25	Glu 65	Pro	Gly	Ala	Leu	Leu 70	Cys	Leu	Thr	Ser	Arg 75	Asp	Gly	Ser	Leu	Leu 80
30	Leu	Ser	Cys	Val	Arg 85	Pro	His	His	Val	Ile 90	Lys	Pro	Lys	Gly	Thr 95	Ala
.	Lys	Lys	Lys	Lys 100	-	Lys	Lys	Lys	Lys 105	Lys	Lys	Lys	Lys	Lys 110	Xaa	Xaa
35	Gly	Gly	Xaa 115													
40	(2)	INF		TION SEQU	ENCE	СНА	RACT	ERIS	TICS							
45			(xi)	((B) 1	YPE :	ami : OGY	no a no a lin PTIO	cid ear			: 31	0:			
	Met 1	-	Leu	Pro	Gln 5		Ile	Tyr	Leu	Phe 10		Phe	Cys	Phe	Cys 15	Cys
50	Leu	Ala	Ile	Val		Asn	Ala	Ser	Ile 25		Ile	His	Ile	Gln 30	Val	Ser
. .	Met	Trp	Leu 35	_	Val	Phe	Ile	Ser 40		Gly	Tyr	Leu	His 45		Ser	Arg
55	Ile	Leu 50	_	' His	Asn	Ile	Ile 55		Cys	Leu	Thr	Ser 60		Arg	Ile	Ala
60	Lys 65	_	Phe	Phe	Ile	Val		Ala	Ser	Phe	Thr 75		Pro	Pro	Ala	Met 80

```
Tyr Lys Asp Phe Tyr Phe Ser Ile Ser Leu His Leu Pro Thr Leu Leu
                                          90
 5
     Phe Xaa Xaa Xaa Phe Val Phe Ser Leu Leu Pro Pro
                 100
                                      105
10
      (2) INFORMATION FOR SEQ ID NO: 311:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 65 amino acids
                    (B) TYPE: amino acid
15
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 311:
     Met Cys Ser Pro Ser Leu Ser Ser Ser Pro Pro Pro Leu Leu Gln Val
20
      Phe Phe Phe Phe Phe Phe Ser Pro His Trp Ala Ala Lys Val Val Pro
                  20
                                      25
      Gln Trp Lys Xaa Arg His Pro Gln Val Ser Ser Gln Leu Leu Cys
25
      Phe Leu Arg Val Asn Cys Gln Phe Leu Phe Leu Gln Glu Ile Leu Phe
30
     Xaa
       65
35
      (2) INFORMATION FOR SEQ ID NO: 312:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 50 amino acids
                    (B) TYPE: amino acid
40
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 312:
      Met Cys Leu Ser Arg Trp Lys Ile Phe Tyr Thr Leu Leu Ile Leu Phe
                                         10
45
      Xaa Xaa Phe Ser Ile Thr Ser Glu Xaa Glu Thr Phe Tyr Met Ile Ile
      Ile His His Asn Pro Thr Gln Ile Thr Ala Ser Cys Ser Phe Thr Phe
50
                                   40
      Leu Xaa
           50
55
      (2) INFORMATION FOR SEQ ID NO: 313:
             (i) SEQUENCE CHARACTERISTICS:
60
                    (A) LENGTH: 293 amino acids
```

(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 313:

	•	(xi)	SEQU	ENCE	DES	CRIE	MIOITS.	l: SE	χ II	O NO:	313	3:			
5	Met Glu 1	Arg :	Pro	Asp 5	Trp	Glu	Thr	Ala	Ile 10	Gln	Lys	Pro	Leu	Cys 15	Ser
10	Leu Pro	Ala (Gly 20	Ser	Gly	Asn	Ala	Leu 25	Ala	Ala	Ser	Leu	Asn 30	His	Tyr
10	Ala Gly	Tyr :	Xaa	Gln	Val	Thr	Asn 40	Glu	Asp	Leu	Leu	Thr 45	Asn	Cys	Thr
15	Leu Leu 50	Leu	Cys	Arg	Arg	Leu 55	Leu	Ser	Pro	Met	Asn 60	Leu	Leu	Ser	Leu
	His Thr 65	Ala	Ser	Gly	Leu 70	Arg	Leu	Phe	Ser	Val 75	Leu	Ser	Leu	Ala	Trp 80
20	Gly Phe	Ile	Ala	Asp 85	Val	Asp	Leu	Glu	Ser 90	Glu	Lys	Tyr	Arg	Arg 95	Leu
25	Gly Glu	Met	Arg 100	Phe	Thr	Leu	Gly	Thr 105	Phe	Leu	Arg	Leu	Ala 110	Ala	Leu
	Arg Thr	Tyr 115	Arg	Gly	Arg	Leu	Ala 120	Tyr	Leu	Pro	Val	Gly 125	Arg	Val	Gly
30	Ser Lys	Thr	Pro	Ala	Ser	Pro 135	Val	Val	Val	Gln	Gln 140	Gly	Pro	Val	Asp
	Ala His 145	Leu	Val	Pro	Leu 150	Glu	Glu	Pro	Val	Pro 155	Ser	His	Trp	Thr	Val 160
35	Val Pro	Asp	Glu	Asp 165	Phe	Val	Leu	Val	Leu 170	Ala	Leu	Leu	His	Ser 175	His
40	Leu Gly	Ser	Glu 180	Met	Phe	Ala	Ala	Pro 185	Met	Gly	Arg	Cys	Ala 190	Ala	Gly
٠.	Val Met	His 195	Leu	Phe	Tyr	Val	Arg 200	Ala	Gly	Val	Ser	Arg 205	Ala	Met	Leu
45	Leu Arg 210		Phe	Leu	Ala	Met 215	Glu	Lys	Gly	Arg	His 220	Met	Glu	Tyr	Glu
	Cys Pro 225	Tyr	Leu	Val	Туг 230		Pro	Val	Val	Ala 235		Arg	Leu	Glu	Pro 240
50	Lys Asp	Gly	Lys	Gly 245		Phe	Ala	Val	Asp 250	_	Glu	Leu	Met	Val 255	Ser
55	Glu Ala	Val	Gln 260	Gly	Gln	Val	His	Pro 265		Туг	Phe	Trp	Met 270		Ser
	Gly Cys	Val 275	Glu	Pro	Pro	Pro	Ser 280	_	Lys	Pro	Gln	Gln 285		Pro	Pro
60	Pro Glu 290		Pro	Leu											

5	(2)	INFO	DRMAT	'ION	FOR	SEQ	ID N	ю: 3	14:							
3			(i) S	-												
				-	A) LI 3) TY					acid	S					
10			(xi)		ייי ארבונייטי די אורבונייטי					20 TI	n Nico	. 31	1.			
10			(XI)	SEQU	/EIVCE	· Dec	CKLI		N. 31	ינ גיב	J 140	. 31.	٠.			
	Met 1	Pro	Leu	Glu	Gly 5	Phe	Cys	Leu	Val	Leu 10	Asp	Ile	Gly	Phe	Leu 15	Leu
15	Val	Met	Leu	Ile 20	Ser	Leu	Ala	Ser	Glu 25	Cys	Phe	Thr	Thr	Cys 30	Leu	Asp
20	Ser	Phe	Ser 35	Thr	Thr	Glu	Pro	Gly 40	Cys	Lys	Phe	Tyr	Lys 45	Leu	Leu	His
	Ser	Val 50	Ser	Leu	Leu	Asn	Ile 55	Asn	Phe	Asn	Val	Lys 60	Ser	Leu	Leu	Суs
25	Ser 65	His	Ile	Xaa			-									
	(2)	INF	ORMA:	rion	FOR	SEQ	ID I	NO: 3	315:							
30			(i)	SEQUI	ENCE	CHA	RACT	ERIS	TICS	:						
				-	A) L B) T					aci	ds					
25				(D) T	OPOL	OGY:	lin	ear				_			
35	·		(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 31	5:			
	Met 1		Leu	Gln	Leu 5	Ser	GĺĄ	Gln	Tyr	Trp 10	Ile	Ser	Leu	Leu	Val 15	Phe
40	Leu	Ser	Leu	Gln 20	Pro	Phe	Pro	Gln	Ala 25	Ala	Ile	Pro	Cys	Ala 30	Leu	Thr
45	Asp	Val	Gly 35	Gly	Ser	Cys	Val	Ile 40	-	His	Ile	Leu	Leu 45	Asn	Cys	Leu
	Cys	Ile 50	Leu	Phe	Thr	Leu	Thr 55	Ala	Pro	Ser	Leu	Ser 60		Val	Leu	Leu
50	Ile 65		Met	Ser	Leu	Ser 70		Суз	Tyr	Glu	Pro 75	Gly	Ala	Asp	Leu	Ser 80
	Asp	Arg	, Ala	Ala	Thr 85	Gly	Asn	Lys	Lys	Leu 90		Arg	Ser	Thr	Су s 95	Leu
55	Leu	Met	: His	Ser 100	Asn	Lys	Leu	Cys	Xaa 105							

60 (2) INFORMATION FOR SEQ ID NO: 316:

		((i) S	-	() Li		1: 7	l am	ino a		s					
5		((xi)	(1) T	OPOL	CGY:	line	ear	O II	ONO:	: 316	:			
10	Met 1	Trp	Gly	Cys	Ser 5	Gly	Leu	Gly	His	Arg 10	Thr	Val	Ser	Phe	Leu :	Leu
	Leu 1	Leu	Pro	Cys 20	Ser	Phe	Pro	Arg	Pro 25	Cys	Xaa	Leu	Phe	30 Gly	Leu	Ile
15	Pro	Ile	Ser 35	Arg	Pro	Cys	Lys	Val 40	Glu	Ala	Pro	Arg	Leu 45	Ser	Val	Pro
	Xaa	Leu 50	Ser	Cys	Ala	Ser	His 55	Pro	Tyr	Cys	Asn	Cys 60	Pro	Met	Ser	Thr
20	Ser 65	Cys	Pro	Leu	Pro	Arg 70	Xaa									
25	(2)	INFO	ORMA!	rion	FOR	SEQ	ID I	NO: 3	317:							
30				0	A) L B) T D) T	ENGT YPE : OPOL	H: 3 ami OGY:	9 am no a lin	ino cid ear	acid		: 317	7 :			
26	Met 1	Leu	Asn	Val	Leu 5	Ser	Lys	Val	Gln	Gln 10	Leu	Val	Ser	Xaa	Leu 15	Gly
35	Leu	Val	Thr	Phe 20	Leu	Leu	Asn	His	Ser 25	Ala	Ala	Gly	Gly	Ser 30	Pro	Gln
40	His	Arg	Trp 35	Leu	Leu	Leu	Xaa									
45	(2)	INF		TION SEQU						١•						
50				(A) I B) T D) T	ENGT TYPE :	H: 7 ami OGY:	72 an ino a : lir	nino acid near	acid						
50	Met	Lvs								-): 31 Ser		Leu	Val	Leu
	1	-,-			5	_		,-		10					15	
55	Pro	His	Val	Val 20		Glu	His	Leu	Phe 25	-	His	His	Asn	Pro 30	Arg	His
60	Pro	Val	11e 35	_	Pro	Phe	Pro	Pro 40		His	Leu	Ile	Ser 45	-	Ser	Val _.

•	Ser	50	Ser	Thr	Trp	HIS	55	GIÀ	GIu	Xaa	Leu	60	Leu	Leu	Val	Pro
5	Ile 65	Ala	Pro	Ser	Val	Trp 70	Ser	Xaa								
10	(2)		ORMAT	SEQUI	ENCE	CHA	RACTI	ERIST	rics		_					
15			(xi)	(1 (1	A) L B) T D) T UENCI	YPE: OPOL	ami OGY:	no a	cid ear			: 319	9:			
	Met 1	Glu	Gln	Gly	Gly 5	Gly	Pro	Arg	Leu	Leu 10	Leu	Leu	Ile	Pro	Gly 15	Leu
20	Leu	His	Asn	Thr 20	Туг	Leu	Ala	Arg	Pro 25	Gly	Asp	Phe	Pro	Ala 30	Gln	Gly
25	Thr	Thr	Glu 35	Asn	Thr	Glu	Cys	Gln 40	Gly	Ser	Pro	Ser	Pro 45	Ile	Ser	His
	Leu	Gly 50	Lys	Val	Arg	Ser	Leu 55	Asp	Ser	Asn	Thr	Gln 60	Ile	Xaa		
30	(2)	INF	ORMA:	rion	FOR	SEQ	ID 1	NO: 3	320:							
35				(ENCE A) L B) T D) T UENC	ENGT YPE: OPOL	H: 2 ami OGY:	86 a no a lin	mino cid ear	aci		. 32	n -			
40	Met 1	Pro	Leu				•							Gly	Ser 15	Val
	Thr	Leu	Gln	Gln 20	Arg	Gly	Met	Phe	Leu 25	Pro	Trp	Thr	Gly	Thr 30	Gly	Glu
45	Gln	Val	Leu 35	Ala	Leu	Leu	Trp	Pro 40	Arg	Phe	Glu	Leu	Ile 45	Leu	Glu	Met
50	Asn	Val 50	Glņ	Ser	Vạl	Arg	Ser 55	Thr	Asp	Pro	Gln	Arg 60	Leu	Gly	Gly	Leu
	Asp 65		Arg	Pro	His	Tyr 70	Ile	Thr	Arg	Arg	Tyr 75	Ala	Glu	Phe	Ser	Ser 80
55	Ala	Leu	Val	Ser	Ile 85	Asn	Gln	Thr	Ile	Pro 90	Asn	Glu	Arg	Thr	Met 95	Gln
			Gly	100					105					110	_	
60	Ala	Ala	Glu	Phe	Ser	Ser	Arg	Lys	Glu	Gln	Leu	Val	Phe	Leu	Ile	Asn

•		115					120					125			
5	Asn Ty 13	-	Met	Met	Leu	Gly 135	Val	Leu	Met	Glu	Arg 140	Ala	Ala	Asp	Asp
J	Ser Ly 145	s Glu	Val	Glu	Ser 150	Phe	Gln	Gln	Leu	Leu 155	Asn	Ala	Arg	Thr	Gln 160
10	Glu Ph	e Ile	Glu	Glu 165	Leu	Leu	Ser	Pro	Pro 170	Phe	Gly	Gly	Leu	Val 175	Ala
	Phe Va	ıl Lys	Glu 180	Ala	Glu	Ala	Leu	Ile 185	Glu	Arg	Gly	Gln	Ala 190	Glu	Arg
15	Leu Ar	g Gly 195		Glu	Ala	Arg	Val 200	Thr	Gln	Leu	Ile	Arg 205	Gly	Phe	Gly
20	Ser Se	_	Lys	Ser	Ser	Val 215	Glu	Ser	Leu	Ser	Gln 220	Asp	Val	Met	Arg
	Ser Ph 225	ne Thr	Asn	Phe	Arg 230	Asn	Gly	Thr	Ser	Ile 235	Ile	Gln	Gly	Ala	Leu 240
25	Thr G	ln Leu	Ile	Gln 245	Leu	Tyr	His	Arg	Phe 250	His	Arg	Val	Leu	Ser 255	Gln
	Pro G	ln Leu	Arg 260		Leu	Pro	Ala	Arg 265		Glu	Leu	Ile	Asn 270	Ile	His
30	His L	eu Met 275		Glu	Leu	Lys	Lys 280		Lys	Pro	Asn	Phe 285	Xaa		
35	(2) II	NFORMA	TION	FOR	SEQ	ID :	NO:	321:							
40			1	(A) I (B) I (D) I	ENGI YPE :	H: S ami OGY:	5 an ino a	nino ncid near	ació): 32	1:			
45	Met P	he Arg	, Ala	Leu 5		Asp	Leu	Leu	Thr 10		Туг	Pro	Gln	Gln 15	
	Leu L	eu Glr	1 Val 20		Val	Val	Met	Tyr 25		Val	Leu	Gln	Val 30		Glu
50	Leu P	ro Try 35		Glu	Leu	Ile	His 40		Gln	Gly	Ile	• Val 45		Thr	Asp
		eu His 50	. Leu	Lys	Gln	Хаа 55									
55															
	(2) I	NFORM													
60		(i)	SEQU	JENCE (A)						ds.					

				(1	T (O	OPOL	OGY:	lin	ear							
			(xi)	SEQU	JENCI	E DES	SCRI	PTIO	1: SI	EQ II	ОИС	: 32	2 :			
5	Asp 1	Phe	Val	Pro	Val 5	Leu	Val	Phe	Val	Leu 10	Ile	Lys	Ala	Asn	Pro 15	Pro
10	Cys	Leu	Leu	Ser 20	Thr	Val	Gln	Tyr	Ile 25	Ser	Ser	Phe	Tyr	Ala 30	Ser	Cys
10	Leu	Ser	Gly 35	Glu	Glu	Ser	Tyr	Trp 40	Trp	Met	Gln	Phe	Thr 45	Ala	Ala	Val
15	Glu	Phe 50	Ile	Lys	Thr	Ile	Asp 55	Asp	Arg	Lys	Xaa					
20	(2)	INF	ORMA					VO: 3		•						
				(A) L B) T	ENGT YPE :	H: 1 ami	20 a no a lin	mino cid		ds					
25			(xi)	SEQ	UENC	E DE	SCRI	PTIO	V: S	EQ I	D NO	: 32	3:			
	Met 1	His	Pro	Ala	Arg 5	Lys	Leu	Leu	Ser	Leu 10	Leu	Phe	Leu	Ile	Leu 15	Met
30	Gly	Thr	Glu	Leu 20	Thr	Gln	Asp	Ser	Ala 25	Ala	Pro	Asp	Ser	Leu 30	Leu	Arg
35	Ser	Ser	Lys 35	Gly	Ser	Thr	Arg	Gly 40	Ser	Leu	Ala	Ala	Ile 45	Val	Ile	Trp
	Arg	Gly 50	Lys	Ser	Glu	Ser	Arg 55		Ala	Lys	Thr	Pro 60	Gly	Ile	Phe	Arg
40	Gly 65		Gly	Thr	Leu	Val 70	Leu	Pro	Pro	Thr	His 75	Thr	Pro	Glu	Trp	Leu 80
•	Ile	Leu	Pro	Leu	Gly 85	Ile	Thr	Leu	Pro	Leu 90	Gly	Ala	Pro	Glu	Thr 95	Gly
45	Gly	Gly	Asp	Cys 100	Ala	Ala	Glu	Thr	Trp 105	Lys	Gly	Ser	Gln	Arg 110	Ala	Gly
50	Gln	Leu	Cys 115	Ala	Leu	Leu	Ala	Xaa 120								
	(2)	INF	ORMA	TION	FOR	SEQ	ID 1	NO:	324:							
55			(i)	(A) I B) I	ENGI YPE :	H: 4	ERIS 4 am no a lin	ino cid		ls					
60			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 32	4:			

530

Phe Phe Leu Val Val Phe Ser Leu Ser Phe Xaa Pro Ser Val Leu Thr 10 Ser Pro Val His Xaa Pro His Cys Cys Gln Xaa Asp Xaa Ile Leu Phe 5 Lys Asn Thr Leu Xaa Xaa Phe Xaa Ala Lys Tyr Xaa 40 35 10 (2) INFORMATION FOR SEQ ID NO: 325: (i) SEQUENCE CHARACTERISTICS: 15 (A) LENGTH: 59 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 325: 20 Met Phe Ser Arg Thr Ser Asn Phe Trp Thr Phe Phe Gln Phe Leu Ile Phe Lys Val Phe Leu Val Leu Lys Asn Xaa Phe Thr Ser Gln Lys 25 Ile Xaa Xaa Ile Xaa Xaa Glu Lys Pro Lys Lys Lys Xaa Arg Gly Gly Arg Ala Pro Ser Pro Gln Gly Gly Pro Xaa 30 50 55 (2) INFORMATION FOR SEQ ID NO: 326: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 326: Met Gly Leu Leu Ile Phe Met Leu Leu Ile Gly Ile His Ser Gln Cys 1 5 10 45 Ser Xaa 50 (2) INFORMATION FOR SEQ ID NO: 327: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 87 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 327: Met Val Leu Phe Cys Phe Val Leu Phe Cys Phe Val Phe Glu Met Asp. 5 60

•	Ser	Ser	Ser	Val 20	Thr	Gln	Ala	Gly	Val 25	Gln	Trp	Суз	Asp	Leu 30	Gly	Ser
5	Leu	Gln	Ala 35	Pro	Pro	Pro	Gly	Phe 40	Ser	Pro	Phe	Ser	Суз 45	Leu	Ser	Leu
	Pro	Ser 50	Ser	Trp	Asp	Tyr	Arg 55	Arg	Pro	Pro	Pro	Arg 60	Pro	Ala	Asn	Phe
10	Leu 65	-	Phe	Leu	Val	Glu 70	Thr	Gly	Phe	His	His 75	Val	Ser	Gln	Asp	Gly 80
15	Leu	Asp	Leu	Leu	Thr 85	Ser	Xaa									
20	(2)	INF	ORMA	UQG2)	ENCE	CHA ENGT	RACT H: 5	ERIS	TICS mino		ds					
25			(xi)	SEQ				lin PTIO		EQ I	D NO	: 32	8 :			
25	Met 1		Thr	Lys	Lys 5		Cys	Ile	Val	Gly 10	Gly	Ile	Leu	Leu	Val 15	Phe
30	Gln	Ile	lle	Ala 20		Leu	Val	Gly	Gly 25	Leu	Ile	Ala	Pro	Gly 30	Pro	Thr
	Thr	Ala	a Val 35		Tyr	Met	Ser	Val 40	Lys	Cys	Val	Asp	Ala 45	Arg	Lys	Asn
35	His	His 50	S Lys)	Thr	Lys	Trp	Phe 55		Pro	Trp	Gly	Pro 60	Asn	His	Cys	Asp
40	Lys 69		e Arg	Asp	Ile	Glu 70		Ala	Ile	Pro	Arg 75		Ile	Glu	Ala	Asn 80
٠.	Asp) Ile	e Val	. Phe	Ser 85		His	Ile	Pro	Leu 90		His	Met	Glu	Met 95	
45	Pro	Tr	p Ph∈	Gln 100		Met	Leu	Phe	Ile 105		Gln	Leu	Asp	Ile 110		Phe
	Lys	s Le	u Asr 115		Glm	lle	Arg	120		Ala	Glu	Val	Ser 125		Asp	Val
50	Sei	13°		Туг	Arg	Asp	Asp 139		Phe	Ala	Glu	140		Glu	Met	Ala
55	Hi:		u Arg	y Val	Pro	Arg 150		: Leu	Lys	Суз	155		Thr	Ser	Pro	Lys 160
	Th	r Pr	o Glu	ı His	165	_	/ Arg	туг	Тух	Glu 170	-	Asp	Val	Leu	175	Phe
60	Met	t Gl	u Ile	Gly 180		· Val	l Ala	A His	Lys 185		тут	Leu	Leu	Asn 190		Arg

	Leu	Pro	Val 195	Asn	Glu	Lys	Lys	Lys 200	Ile	Asn	Val	Gly	Ile 205	Gly	Glu	Ile
5	Lys	Asp 210	Ile	Arg	Leu	Val	Gly 215	Ile	His	Gln	Asn	Gly 220	Gly	Phe	Thr	Lys
10	Val 225	Trp	Phe	Ala	Met	Lys 230	Thr	Phe	Leu	Thr	Pro 235	Ser	Ile	Phe	Ile	Ile 240
	Met	Val	Trp	Tyr	Trp 245	Arg	Arg	Ile	Thr	Met 250	Met	Ser	Arg	Pro	Pro 255	Val
15	Leu	Leu	Glu	Lys 260	Val	Ile	Phe	Ala	Leu 265	Gly	Ile	Ser	Met	Thr 270	Phe	Ile
	Asn	Ile	Pro 275	Val	Glu	Trp	Phe	Ser 280	Ile	Gly	Phe	Asp	Trp 285	Thr	Trp	Met
20		290					295	Gln				300				
2 5	305					310		Gly			315					320
					325			Trp		330					335	
30				340				Phe	345					350		
			355					360					365			Glu
35		370)				375					380				Tyr
40	385					390					395					Ser 400
					405					410)				415	
45				420)				425	•				430		Leu
			435	5				440	,				445	•		Thr
50		450					455	5				460)			Ala
55	465	5				470)				479	5				480
					485	5				490)				495	
60	Asr	ı Gl	y Met	Gl: 500		ı Pro	Cys	s Lys	Ser 509		g Gl	ı Ası	Cys	510		ı Phe

533

•	Val	Ser	Glu 515	Leu	Tyr	Gln	Glu	Leu 520	Phe	Ser	Ala	Ser	Lys 525	Tyr	Ser	Phe
5	Ile	Asn 530	Asp	Asn	Ala	Ala	Ser 535	Gly	Ile	Xaa						
10	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	ю: 3	329:							
15				(A) L B) T D) T	ENGT YPE: OPOL	RACTI H: 2 ami OGY: SCRI	02 a no a lin	mino cid ear	aci		: 32	9:			
20	Met 1	Gly	Ile	Ala	Leu 5	Ala	Val	Leu	Gly	Trp 10	Leu	Ala	Val	Met	Leu 15	Cys
20	Cys	Ala	Leu	Pro 20	Met	Trp	Arg	Val	Thr 25	Ala	Phe	Ile	Gly	Ser 30	Asn	Ile
25	Val	Thr	Ser 35	Gln	Thr	Ile	Trp	Glu 40	Gly	Leu	Trp	Met	Asn 45	Cys	Val	Val
	Gln	Ser 50	Thr	Gly	Gln	Met	Gln 55	Cys	Lys	Val	Tyr	Asp 60	Ser	Leu	Leu	Ala
30	Leu 65	Pro	Gln	Asp	Leu	Gln 70	Ala	Ala	Arg	Ala	Leu 75	Val	Ile	Ile	Ser	Ile 80
35	Ile	Val	Ala	Ala	Leu 85	Gly	Val	Leu	Leu	Ser 90	Val	Val	Gly	Gly	Lys 95	Cys
33	Thr	Asn	Cys	Leu 100	Glu	Asp	Glu	Ser	Ala 105	Lys	Ala	Lys	Thr	Met 110	Ile	Val
40	Ala	Gly	Val 115	Val	Phe	Leu	Leu	Ala 120	Gly	Leu	Met	Val	Ile 125	Val	Pro	Val
٠.	Ser	Trp 130	Thr	Ala	His	Asn	Ile 135	Ile	Gln	Asp	Phe	Tyr 140	Asn	Pro	Leu	Val
45	Ala 145	Ser	Gly	Gln	Lys	Arg 150	Glu			Ala			Tyr		Gly	
50	Ala	Ala	Ser	Gly	Leu 165	Leu	Leu	Leu	Gly	Gly 170	Gly	Leu	Leu	Cys	Cys 175	Asn
50	Cys	Pro	Pro	Arg 180	Thr	Asp	Lys	Pro	Туг 185	Ser	Ala	Lys	Tyr	Ser 190	Ala	Ala
55	Arg	Ser	Ala 195	Ala	Ala	Ser	Asn	Tyr 200	Val	Xaa						

(2) INFORMATION FOR SEQ ID NO: 330:

			(i) :	-	ENCE A) L						đe					
					B) T					acı	us					
5			(xi)		D) IN JENCI					eo II	O NO	: 330	3 :			
,			(**)	320	J	, ,,,							•			
	Met 1	Ala	Thr	Val	Thr 5	Ala	Thr	Thr	Lys	Val 10	Pro	Glu	Ile	Arg	Asp 15	Val
10	Thr	Arg	Ile	Glu 20	Arg	Ile	Gly	Ala	His 25	Ser	His	Ile	Arg	Gly 30	Leu	Gly
15	Leu	Asp	Asp 35	Ala	Leu	Glu	Pro	Arg 40	Gln	Ala	Ser	Gln	Gly 45	Met	Val	Gly
13	Gln	Leu 50	Ala	Ala	Arg	Arg	Ala 55	Ala	Gly	Val	Val	Leu 60	Glu	Met	Ile	Arg
20	Glu 65	Gly	Lys	Ile	Ala	Gly 70	Arg	Ala	Val	Leu	Ile 75	Ala	Gly	Gln	Pro	Gly 80
	Thr	Gly	Lys	Thr	Ala 85	Ile	Ala	Met	Gly	Met 90	Ala	Gln	Ala	Leu	Gly 95	Pro
25	Asp	Thr	Pro	Phe 100	Thr	Ala	Ile	Ala	Gly 105	Ser	Glu	Ile	Phe	Ser 110	Leu	Glu
30	Met	Ser	Lys 115		Glu	Ala	Leu	Thr 120	Gln	Ala	Phe	Arg	Arg 125	Ser	Ile	Gly
	Val	Arg 130		Lys	Glu	Glu	Thr 135	Glu	Ile	Ile	Glu	Gly 140	Glu	Val	Val	Glu
35	Ile 145		Ile	Asp	Arg	Pro 150	Ala	Thr	Gly	Thr	Gly 155	Ser	Lys	Val	Gly	Lys 160
	Leu	Thr	Leu	Lys	Thr 165	Thr	Glu	Met	Glu	Thr 170		Tyr	Asp	Leu	Gly 175	Thr
40	Lys	Met	Ile	Xaa 180	Ser	Leu	Thr	Lys	Asp 185		Val	Gln	Ala	Gly 190		Val
45	Ile	Thr	Ile 195	_	Lys	Ala	Thr	Gly 200		Ile	Ser	Lys	Leu 205		Arg	Ser
	Phe	Thr 210		Ala	Arg	Glu	Leu 215		Arg	Tyr	Gly	Leu 220		Ąsp	Gln	Val
50	Arg 225		Val	Pro	Arg	Trp 230	_	Ala	Pro	Glu	Thr 235		Gly	Gly	Gly	Ala 240
	His	Arg	Val	Pro	Ala 245		Asp	Arg	Arg	His 250		Leu	Ser	His	Pro 255	
55	Leu	Pro	Gly	Ala 260	Leu	Leu	Arg									

60 (2) INFORMATION FOR SEQ ID NO: 331:

5				(A) I B) T D) T	ENGT YPE : YPOL	H: 2 ami OGY:	60 a no a lin	mino cid ear	aci		: 33	1.			
10	Met 1	Leu		Leu										Leu	Leu 15	Gly
10	Leu	Lys	Gly	Leu 20	Ala	Pro	Ala	Glu	Ile 25	Ser	Ala	Val	Cys	Glu 30	Lys	Gly
15	Asn	Phe	Asn 35	Val	Ala	His	Gly	Leu 40	Ala	Trp	Ser	Tyr	Tyr 45	Ile	Gly	Туг
	Leu	Arg 50	Leu	Ile	Leu	Pro	Glu 55	Leu	Gln	Ala	Arg	Ile 60	Arg	Thr	Tyr	Asn
20	Gln 65	His	Tyr	Asn	Asn	Leu 70	Leu	Arg	Gly	Ala	Val 75	Ser	Gln	Arg	Leu	Tyr 80
25	Ile	Leu	Leu	Pro	Leu 85	Asp	Cys	Gly	Val	Pro 90	Asp	Asn	Leu	Ser	Met 95	Ala
	Asp	Pro	Asn	Ile 100	Arg	Phe	Leu	Asp	Lys 105	Leu	Pro	Gln	Gln	Thr 110	Gly	Asp
30	Arg	Ala	Gly 115	Ile	Lys	Asp	Arg	Val 120	Tyr	Ser	Asn	Ser	Ile 125	Tyr	Glu	Leu
	Leu	Glu 130	Asn	Gly	Gln	Arg	Ala 135	Gly	Thr	Суѕ	Val	Leu 140	Glu	Tyr	Ala	Thr
35	Pro 145	Leu	Gln	Thr	Leu	Phe 150	Ala	Met	Ser	Gln	Туг 155	Ser	Gln	Ala	Gly	Phe 160
10	Ser	Gly	Glu	Asp	Arg 165	Leu	Gľu	Gln	Ala	Lys 170	Leu	Phe	Cys	Arg	Thr 175	Leu
	Glu	Asp	Ile	Leu 180	Ala	Asp	Ala	Pro	Glu 185	Ser	Gln	Asn	Asn	Cys 190	Arg	Leu
1 5	Ile	Ala	Туг 195	Gln	Glu	Pro	Ala	Asp 200	Asp	Ser	Ser	Phe	Ser 205	Leu	Ser	Gln
	Glu	Val 210	Leu	Arg	His	Leu	Arg 215	Gln	Glu	Glu	Lys	Glu 220	Glu	Val	Thr	Val
50	Gly 225	Ser	Leu	Lys	Thr	Ser 230	Ala	Val	Pro	Ser	Thr 235	Ser	Thr	Met	Ser	Gln 240
55	Glu	Pro	Glu	Leu	Leu 245	Ile	Ser	Gly	Met	Glu 250	Lys	Pro	Leu	Pro	Leu 255	Arg
	Thr	Ąsp	Phe	Ser 260												

536

(2) INFORMATION FOR SEQ ID NO: 332:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 48 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 332:

Met Thr Pro Gln Lys Pro Ala Leu Ala Val Leu Leu Leu Glu Val Pro 10 1 5 10 15

Leu Leu Leu Thr Leu Ser Val Leu Lys Lys Arg Cys Leu Val Thr Cys 20 25 30

Glu Pro Thr Ser Arg Phe Val Ser Cys Asp Leu Pro Leu Ser Val Xaa 35 40 45

20

30

60

5

(2) INFORMATION FOR SEQ ID NO: 333:

25 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 334 amino acids

- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 333:

Met Ala Ala Ala Trp Leu Gln Val Leu Pro Val Ile Leu Leu Leu

Leu Gly Ala His Pro Ser Pro Leu Ser Phe Phe Ser Ala Gly Pro Ala 35 20 25 30

Thr Val Ala Ala Asp Arg Ser Lys Trp His Ile Pro Ile Pro Ser

40 Gly Lys Asn Tyr Phe Ser Phe Gly Lys Ile Leu Phe Arg Asn Thr Thr 50 55 60

Ile Phe Leu Lys Phe Asp Gly Glu Pro Cys Asp Leu Ser Leu Asn Ile 65 70 75 80

Thr Trp Tyr Leu Lys Ser Ala Asp Cys Tyr Asn Glu Ile Tyr Asn Phe 85 90 95

Lys Ala Glu Glu Val Glu Leu Tyr Leu Glu Lys Leu Lys Glu Lys Arg 50 100 105 110

Gly Leu Ser Gly Lys Tyr Gln Thr Ser Ser Lys Leu Phe Gln Asn Cys 115 120 125

55 Ser Glu Leu Phe Lys Thr Gln Thr Phe Ser Gly Asp Phe Met His Arg

Leu Pro Leu Gly Glu Lys Gln Glu Ala Lys Glu Asn Gly Thr Asn 145 150 155 160

	Leu	Thr	Phe	Ile	Gly 165	Asp	Lys	Thr	Ala	Met 170	His	Glu	Pro	Leu	Gln 175	Thr	
5	Trp	Gln	Asp	Ala 180	Pro	Tyr	Ile	Phe	Ile 185	Val	His	Ile	Gly	Ile 190	Ser	Ser	•
	Ser	Lys	Glu 195	Ser	Ser	Lys	Glu	Asn 200	Ser	Leu	Ser	Asn	Leu 205	Phe	Thr	Met	=
0	Thr	Val 210	Glu	Val	Lys	Gly	Pro 215		Glu	Tyr	Leu	Thr 220		Glu	Asp	Туз	•
15	Pro 225		Met	Ile	Phe	Phe 230		Val	Met	Cys	Ile 235		Tyr	Val	Leu	240	e 0
			Leu		245					250					255	•	
20			Gln	260					265	ı				270)		
			Val 275	i				280	}				285	•			
25		290					295	5				300)				
30	305	5	r Leu			310)				319	5				r G1 32	.Y
	Ile	e Vai	l Lys	s Pro	329		ı Glu	ı Sei	c Le	33(e Ar	g Le	u Xa	a		
35	(2) IN	FORM	ATIO	N FOI	R SE	O ID	NO:	334	:							
40			(i)	SEQ	(A) (B)	LENG TYPE	TH:	TERI 200 nino : li	amin acid	o ac I	ids						
			(xi) SE	QUEN	CE D	ESCR	IPTI	ON:	SEQ	ID N	10: 3	34:				
45		t Va 1	l Le	u Xa		1 V a 5	l Th	r Le	u Gl		u Al .0	a Le	u Ph	e Th		eu C .5	ys
	G1	у Гу	rs Ph		s Ar O	g Tr	p Ly	s Le		n G1 5	lA ų.	a Ph	e Le	eu Le	u I] 10	le T	hr
50	A]	ia Ph	ne Le 3	eu Se 15	er Va	l Le	u Il		p Va 10	נא נו	la Tr	np Me		r Me 15	et Ty	/r L	eu
5 5	Pł		ly As 50	in Va	ıl Ly	rs Le		ln G1 55	ln G]	y As	sp Al		np As 50	en As	sp P	ro T	hr
JJ		eu A] 65	la II	le Tì	ır Le		la Al 70	la Se	er Al	la G		er Se 75	er S	er S	er S	er 1	hr 80
60	P	ro S	er Le	eu Ai		er Ti	nr A	la Pi	ro Pl		ys G: 90	ln P	ro C	ys A	rg A	rg 1 95	lhr

	Arg	Pro	Thr	Thr 100	Ser	Thr	Arg	Arg	Ser 105	Pro	Gly	Cys	Gly	Arg 110	Arg	Pro
5	Ser	Arg	Arg 115	Thr	Суѕ	Ser	Суз	Arg 120	Gly	Pro	Ile	Trp	Arg 125	Thr	Arg	Pro
10	Ser	Pro 130	Trp	Met	Asn	Thr	Met 135	Gln	Leu	Ser	Glu	Gln 140	Gln	Asp	Phe	Pro
10	Thr 145	Ala	Ala	Trp	Glu	Lys 150	Asp	Pro	Val	Ala	Ala 155	Trp	Gly	Lys	Asp	Pro 160
15	Ala	Leu	Arg	Leu	Glu 165	Ala	Thr	Cys	Ile	Ser 170	Gln	Leu	Arg	Trp	Pro 175	Ser
	Суз	Ser	Thr	Val 180	Gly	Pro	Ser	Gln	Leu 185	Leu	Arg	Gln	Val	Thr 190	Gln	Glu
20	Хаа	Thr	Phe 195	Gly	Glu	Arg	Leu	Хаа 200					-			
25	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: :	335:							
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids (B) TYPE: amino acid															
30			(xi)	(D) T	OPOL	OGY:	lin	ear	eo i	D NO	: 33	5:			
35	Met 1	Leu	Leu	His	His 5	Gln	Leu	Leu	Ile	Val 10	Thr	Leu	His	Leu	Val 15	Leu
	Leu	Leu	Ala	Thr 20	Leu	Leu	Val	Xaa								
40	(2)	INF	ORMA:	TION	FOR	SEQ	ID 1	NO:	336:							
45				(ENCE (A) I (B) I (D) I	ENGI YPE : OPOI	H: 1 ami OGY:	.43 a no a lin	mino cid ear	aci		o: 33	6:			
50	Met 1		Lys	Ala	Leu 5		Ile	Tyr	Leu	Val 10	Ser	Ser	Phe	Leu	Ala 15	Leu
	Asn	Gln	Ala	Ser 20		Ile	Ser	Arg	Cys 25	Asp	Leu	Ala	Gln	Val 30	Leu	Gln
55	Leu	Glu	Asp 35		Asp	Gly	Phe	Glu 40	Gly	Tyr	Ser	Leu			Trp	Leu
			33										45			

•	Asn 65	Ala	Asp	Gly	Ser	Phe 70	Asp	Тук	Gly	Leu	Phe 75	Gln	Ile	Asn	Ser	His 80
5	Tyr	Trp	Cys	Asn	Xaa 85	Tyr	Lys	Ser	Tyr	Ser 90	Glu	Asn	Leu	Cys	His 95	Val
	Asp	Суз	Gln	Asp 100	Leu	Leu	Asn	Pro	Asn 105	Leu	Leu	Ala	Gly	Ile 110	His	Cys
10	Ala	Lys	Arg 115	Ile	Val	Ser	Gly	Ala 120	Arg	Gly	Met	Asn	Asn 125	Trp	Val	Arg
15	Met	Glu 130	Xaa	Cys	Thr	Val	Gln 135	Ala	Gly	His	Ser	Ser 140	Thr	Gly	Xaa	
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: 3	337:							
20			(i)	(ENCE A) L B) T D) T	engt Ype:	H: 9 ami	5 am no a	ino cid		s					
25			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 33	7:			
	Met 1		Val	Ile	Ala 5	Gly	Gly	Ile	Leu	Ala 10	Ala	Leu	Leu	Leu	Leu 15	Ile
30	Val	Val	Val	Leu 20	Cys	Leu	Tyr	Phe	Lys 25	Ile	His	Asn	Ala	Leu 30	Lys	Ala
	Ala	Lys	Glu 35	Pro	Glu	Ala	Val	Ala 40	Val	Lys	Asn	His	Asn 45	Pro	Asp	Lys
35	Val	Trp 50		Ala	Lys	Asn	Ser 55		Ala	Lys	Thr	Ile 60	Ala	Thr	Glu	Ser
40	Cys 65		Ala	Leu	Gln	Cys 70	Cys	Glu	Gly	Тух	Arg 75	Met	Cys	Ala	Ser	Phe 80
	Asp	Ser	Leu	Pro	Pro 85	Cys	Cys	Cys	Asp	Ile 90	Asn	Glu	Gly	Leu	Xaa 95	
45	(2)	INF	ORMA	TION	FOR	SEQ	ID 1	NO: I	338:							
			(i)	_	ENCE						~					
50			(xi)	(B) I D) I	YPE: OPOL	ami OGY:	no a lin	cid ear			: 33	8:			
55	Met 1		Leu	Lys	Ser 5	Asn	Ile	Leu	Met	Leu 10	Asn	Leu	Phe	Ala	Ala 15	Asn
	Val	Gly	Ala	Asn 20	Phe	Ala	Leu	Thr	Val 25	Glu	Lys	Ile	Gly	Met 30	Ile	Leu
60	Leu	Asn	Val	Ser	Gly	Xaa										

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540

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5
      (2) INFORMATION FOR SEQ ID NO: 339:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 39 amino acids
                    (B) TYPE: amino acid
10
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 339:
      Met Leu Val Val Ala Phe Gly Leu Leu Val Leu Tyr Ile Leu Leu Ala
15
      Ser Ser Trp Lys Arg Pro Glu Pro Gly Ile Leu Thr Asp Arg Gln Pro
                   20
                                       25
                                                           30
      Leu Leu His Asp Gly Glu Xaa
20
              35
      (2) INFORMATION FOR SEQ ID NO: 340:
25
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 71 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 340:
      Ser Asp Pro Leu Ala Ser Ala Ser Gln Asn Ala Gly Ile Val Ser Val
                                          10
35
      Gly Leu Cys Thr Arg Pro Gly Pro Gln Phe Lys Asn Ala Gln Pro Pro
                  20
                                      25
      Phe Pro Xaa Gln Lys Ala Pro Arg Cys Leu Trp Glu Asn Gln Pro Pro
40
      Pro Trp Arg Lys Ala Trp Asp Leu Pro Ser His Leu Gly Arg Arg Gly
      Ile Cys Gly Lys Ser Phe Xaa
45
      (2) INFORMATION FOR SEQ ID NO: 341:
50
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 85 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
55
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 341:
      Tyr Val Met Ile Phe Lys Lys Glu Phe Ala Pro Ser Asp Glu Glu Leu
       1
                        5
                                           10
60
     Asp Ser Tyr Arg Arg Gly Glu Glu Trp Asp Pro Gln Lys Ala Glu Glu
```

	20	25	30
5	Lys Arg Asn Xaa Lys Glo 35	ı Leu Ala Gln Arg Gln 40	Xaa Gly Gly Gly Ser 45
J	Pro Ala Gly Ala Cys Gly 50	Gly Glu Pro Cys Gln 55	Arg Leu Gln Gly Gln 60
10	Val Gln Pro Pro His Are	-	Arg Arg Ser Pro His
	Ala Thr Gly Gln Xaa 85		•
15			
	(2) INFORMATION FOR SE	Q ID NO: 342:	
20	(i) SEQUENCE CH (A) LENG	ARACTERISTICS: TH: 90 amino acids	
	(B) TYPE	: amino acid	
		LOGY: linear ESCRIPTION: SEQ ID NO:	: 342:
25	Met Trp Asp Trp Asp Tr	n Ser Ala Pro Trn Ser	Trn Pro Leu Trn Leu
	1 5	10	15
20	Ser Leu Ala Leu Val Cy 20	s Leu Ser Ala Gly Ala 25	Lys Gly His Arg Ala 30
30	Ser Glu Ala Gly His Al	a Arg Ala Leu Thr Cys 40	Glu Met Gly Ser Glu 45
35	Phe Xaa Thr Ala Xaa Gl	y Leu Val Leu Gly Xaa 55	Xaa Xaa Trp Thr Xaa 60
	Xaa Asn Gly Ser Ala Gl 65 7	y Pro Glu Arg Arg Gly 0 75	Trp Arg Pro Ala Ala 80
40	Phe Leu Ala Val Phe Le 85	u Leu Gly Asp Xaa 90	
•			
45	(2) INFORMATION FOR SE	Q ID NO: 343:	
		ARACTERISTICS: FTH: 48 amino acids E: amino acid	
50	(D) TOPO	DLOGY: linear DESCRIPTION: SEQ ID NO	: 343:
55	Met Phe Gly Pro Thr Ph	e His Ser Leu Val Leu 10	Val Pro Pro Trp Pro 15
<i>JJ</i>	Asn Leu Ser Leu Leu Hi	s Phe Thr Ser Pro Val 25	Gly Gln His Ser Ser 30
60	Phe Leu Pro Thr Ser Le	u Arg Leu Xaa Lys Lys 40	Lys Lys Lys Lys Lys

5																
	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	10: 3	44:							
10				(1	A) Li B) T	engti YPE: OPOL	H: 5 amin CGY:	6 am no a lin	ino a cid ear	acid		: 34	4:			
15	Met 1	Cys	Ser	Lys	Asn 5	Gly	Phe	Leu	Leu	Ala 10	Trp	Ser	Trp	Asn	Ser 15	Pro
20	Trp	Leu	Pro	Gln 20	Ala	Ser	Leu	Ala	His 25	Gly	Cys	Trp	Gly	Arg 30	Тгр	Met
20	Ser	Asp	Leu 35	Val	Gly	Cys	Ser	Arg 40	Glu	Asn	Lys	Cys	Ala 45	Leu	Arg	Asp
25	His	Ser 50	Glu	Arg	Val	Gln	Gly 55	Xaa								
30	(2)	INF				CHAI	RACT H: 2	ERIS 22 a	TICS mino		ds					
35			(xi)	SEQ	D) T					EQ I	D NO	: 34	5:			
	Ser 1	Pro	Leu	Xaa	Phe 5	Cys	Val-	Val	Leu	Leu 10	Leu	Gln	Ala	Ala	Arg 15	Gly
40	Туг	Val	Val	Arg 20	Lys	Pro	Ala	Gln	Ser 25	Arg	Leu	Asp	Asp	Asp 30	Pro	Pro
45	Pro	Ser	Thr 35	Leu	Leu	Lys	Asp	Туг 40	Gln	Asn	Val	Pro	Gly 45	Ile	Glu	Lys
	Val	Asp 50	Asp	Val	Val	Lys	Arg 55	Leu	Leu	Ser	Leu	Glu 60	Met	Ala	Asn	Lys
50	Lys 65	Glu	Met	Leu	Lys	Ile 70	Lys	Gln	Glu	Gln	Phe 75	Met	Lys	Lys	Ile	Val 80
	Ala	Asn	Pro	Glu	Asp 85	Thr	Arg	Ser	Leu	Glu 90	Ala	Arg	Ile	Ile	Ala 95	Leu
55	Ser	Val	Lys	Ile 100	Arg	Ser	Tyr	Glu	Glu 105	His	Leu	Glu	Lys	His 110	Arg	Lys

Asp Lys Ala His Lys Arg Tyr Leu Leu Met Ser Ile Asp Gln Arg Lys 115 120 125

-	Lys	Met 130	Leu	Lys	Asn	Leu	Arg 135	Asn	Thr	Asn	Tyr	Asp 140	Val	Phe	Glu	Lys
5	Ile 145		Trp	Gly	Leu	Gly 150	Ile	Glu	Tyr	Thr	Phe 155	Pro	Pro	Leu	Tyr	Tyr 160
	Arg	Arg	Ala	His	Arg 165	Arg	Phe	Val	Thr	Lys 170	Lys	Ala	Leu	Cys	11e 175	Arg
10	Val	Phe	Gln	Glu 180	Thr	Gln	Lys	Leu	Lys 185	Lys	Arg	Arg	Arg	Ala 190	Leu	Lys
15	Ala	Ala	Ala 195	Ala	Ala	Gln	Lys	Gln 200	Ala	Lys	Arg	Arg	Asn 205	Pro	Asp	Ser
	Pro	Ala 210	_	Ala	Ile	Pro	Lys 215	Thr	Leu	Lys	Asp	Ser 220	Gln	Xaa		
20	(2)	INF	ORMA	tion	FOR	SEQ	ID I	NO:	346:							
			(i)	SEQU	ENCE	СНА	RACT	ERIS	TICS	:						
25					(A) L (B) T					acid	ls					
			(xi)	((D) I	OPOL	OGY:	lir	ear	EQ I	D NO	: 34	6:			
30	Met 1	_	Ala	Pro	Ala 5		Ser	Leu	Leu	Leu 10	Leu	Leu	Leu	Leu	Phe 15	Ala
	Cys	Cys	Trp	Ala 20		Gly	Gly	Ala	Asn 25	Leu	Ser	Gln	Asp	Asp 30	Ser	Gln
35	Pro	Trp	Thr 35		Asp	Glu	Thr	Val 40		Ala	Gly	Gly	Thr 45		Val	Leu
	Lys	Cys 50		Val	Lys	Asp	His 55		Asp	Ser	Ser	Leu 60		Trp	Ser	Xaa
40																
45																
43	(2)	INE	ORMA	ADITA	FOR	SEC	ID	NO:	347 :							
			(i)	-	JENCI (A)						ids					
50			(xi)		(B) '	TYPE TOPO	: am LOGY	ino a : li	acid near): 34	17 :			
55		t Val	l Ala	a Pro	Val		туг	: Let	val	. Ala		a Ala	. Leu	. Leu	Val	Gly
	Ph	e Ile	e Leu	1 Phe 20		ı Thi	: Arç	g Sei	Arg 25		/ Arg	, Ala	a Ala	Ser 30		a Gly
60	Gl	n Glu	ı Pro	Le	ı His	a Ası	ı Glu	ı Glu	Leu	Ala	Gly	/ Ala	a Gly	/ Arc	y Vai	l Ala

•	35	40	45	
5	Gln Pro Gly Pro Leu (Glu Pro Glu Glu Pr 55	ro Arg Ala Gly 60	Gly Arg Pro
3	Arg Arg Arg Arg Asp 65	Leu Gly Ser Arg Le 70	eu Gln Ala Gln 75	Arg Arg Ala 80
10 ⁻	Gln Arg Val Ala Trp 85	-	lu Asn Glu Glu 90	Glu Ala Val 95
	Ile Leu Ala Gln Glu (Glu Glu Gly Val G 105	lu Lys Pro Ala	Glu Xaa His 110
15	Leu Ser Gly Lys Ile (Gly Ala Lys Lys Le 120	eu Arg Xaa Xaa 125	Glu Glu Lys
20	Gln Ala Arg Lys Ala (Gln Xaa Glu Ala G 135	lu Glu Ala Glu 140	Arg Glu Xaa
	Arg Lys Arg Leu Glu 145	Ser Gln Arg Glu Xa 150	aa.	
25	(2) INFORMATION FOR	SEQ ID NO: 348:		
30	(A) LE (B) TY (D) TC	CHARACTERISTICS: ENGTH: 17 amino ac (PE: amino acid OPOLOGY: linear C DESCRIPTION: SEQ		
35	Met Gln Lys Cys Met 1 5		al Phe His Ile 10	Gln Trp Ser 15
	Xaa			
40			•	
٠.	(2) INFORMATION FOR	SEQ ID NO: 349:		
45	LI (A) TI (B) TI (D)	CHARACTERISTICS: ENGTH: 10 amino ac PE: amino acid DPOLOGY: linear E DESCRIPTION: SEQ		•
50	Met Leu Val Cys Ser 1 5		a a 10	
55	(2) INFORMATION FOR	SEQ ID NO: 350:		
	· · · · · · · · · · · · · · · · · · ·	CHARACTERISTICS: ENGTH: 14 amino ac	:ids	
60		YPE: amino acid OPOLOGY: linear		

```
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 350:
     Val Ile Glu Leu Cys Val Ser Leu Arg Ser Leu Asn Phe Xaa
5
      (2) INFORMATION FOR SEQ ID NO: 351:
10
             (i) SEQUENCE CHARACTERISTICS:
                   (A) LENGTH: 18 amino acids
                    (B) TYPE: amino acid
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 351:
15
     Met Cys Glu Phe Xaa Xaa Xaa Ile Met Xaa Leu Ala Gly Tyr Phe Ala
              5
                              10
     Cys Xaa
20
      (2) INFORMATION FOR SEQ ID NO: 352:
25
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 62 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 352:
      Met Val Gly Gly Tyr Val Ser Ser Phe Ser Phe Pro Pro Val Ser Ser
                                         10
      Ser Leu Leu Pro Ala Ser Phe Ala Phe Pro Phe Leu Pro Gly Thr
35
                                      25
      Pro Cys Pro Phe Leu Tyr Phe Leu Pro Ser Pro Phe Ser Pro Leu Pro
                           40
40
      Leu Ser Leu Thr Arg Ser Asn Ser Phe Leu Leu Asn Gly Xaa
                             55
           50
45
      (2) INFORMATION FOR SEQ ID NO: 353:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amino acids
50
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 353:
      Glu Lys Lys Ser Met Ser Val Ser Asp Ile Tyr Ala Leu Glu Ser Leu
55
      Gly Arg Ser Leu Phe Thr Leu Asn Ser Met Cys Leu Pro Leu Ser Phe
                 20
                                   25
60
```

5	(2)	INFO	ORMAT	ION	FOR	SEQ	ID N	ю: 3	54:							
10			(i) S (xi)	(2 (1 (1	A) LI B) T D) T	ENGTI YPE : OPOLA	d: 2 ami CGY:	45 ar no ac line	mino cid ear	acio		: 354	l:			
15	Met 1	Gly	Gly	Ala	Ser 5	Arg	Arg	Val	Glu	Ser 10	Gly	Ala	Trp	Ala	Tyr 15	Leu
13	Ser	Pro	Leu	Val 20	Leu	Arg	Lys	Glu	Leu 25	Glu	Ser	Leu	Val	Glu 30	Asn	Glu
20	Gly	Ser	Glu 35	Val	Leu	Ala	Leu	Pro 40	Glu	Leu	Pro	Ser	Ala 45	His	Pro	Ile
	Ile	Phe 50	Trp	Asn	Leu	Leu	Trp 55	Tyr	Phe	Gln	Arg	Leu 60	Arg	Leu	Pro	Ser
25	Ile 65	Leu	Pro	Gly	Leu	Val 70	Leu	Ala	Ser	Cys	Asp 75	Gly	Pro	Ser	Xaa	Ser 80
30	Gln	Ala	Pro	Ser	Pro 85	Trp	Leu	Thr	Pro	Asp 90	Pro	Ala	Ser	Val	Gln 95	Val
50	Arg	Leu	Leu	Trp 100	Asp	Val	Leu	Thr	Pro 105	Asp	Pro	Asn	Ser	Cys 110	Pro	Pro
35	Leu	Tyr	Val 115	Leu	Trp	Arg	Val	His 120	Ser	Gln	Ile	Pro	Gln 125	Arg	Val	Val
	Trp	Pro 130		Pro	Val	Pro	Ala 135		Leu	Ser	Leu	Ala 140	Leu	Leu	Glu	Ser
40	Val 145		Arg	His	Val	Gly 150	Leu	Asn	Glu	Val	His 155		Ala	Val	Gly	Leu 160
45	Leu	Leu	Glu	Thr	Leu 165		Pro	Pro	Pro	Thr 170		Leu	His	Leu	Gln 175	Arg
73	Gly	Ile	Tyr	Arg 180		Ile	Leu	Phe	Leu 185		Met	Ala	Ala	Leu 190		Lys
50	Asp	His	Val 195		Ile	Val	Ala	Phe 200		Lys	Lys	Tyr	Lys 205	Ser	Ala	Phe
	Asn	Lys 210		Ala	Ser	Ser	Met 215		Lys	Glu	Glu	Leu 220		His	Arg	Arg
55	Ala 225		n Met	Pro	Thr	230		: Ala	Ile	Asp	Cys 235		Lys	Cys	Phe	Gly 240
60	Ala	Pro	Pro	Glu	Cys 245											

	(2) INFORMITON FOR SEQ ID NO: 555:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 35 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 355:
10	Met Lys Phe Ser Leu Leu Phe Leu Pro Met Leu Leu Ile Leu Lys Pro 1 5 10 15
15	Asp Leu Phe His Ile Ser Ile Cys Thr Leu Ala Ala Cys Gly Leu Thr 20 25 30
	Phe Pro Xaa 35
20	
	(2) INFORMATION FOR SEQ ID NO: 356:
25	(i) SEQUENCE CHARACTERISTICS:
25	(A) LENGTH: 22 amino acids
	(B) TYPE: amino acid (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 356:
20	
30	Met Leu Phe Phe Phe Ile Leu His Leu Leu Ser Ile Met Ser Phe Leu 1 5 10 15
	Ser Pro Asp Ile Met Xaa
25	20
35	
	(2) INFORMATION FOR SEQ ID NO: 357:
40	
40	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 98 amino acids (B) TYPE: amino acid
• •	(D) TOPOLOGY: linear
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 357:
45	Mot Dho Cly I ou tou Wal Cly Con Cla diby I ou Tou Cly Cly Arm Ala
	Met Phe Gly Leu Leu Val Glu Ser Gln Thr Leu Leu Glu Glu Asn Ala 1 5 10 15
50	Val Gln Gly Thr Glu Arg Thr Leu Gly Leu Asn Ile Ala Pro Phe Ile
50	20 25 30
	Asn Gln Phe Gln Val Pro Ile Arg Val Phe Leu Asp Leu Ser Ser Leu
	35 40 45
e =	
55	Pro Cys Ile Pro Leu Ser Lys Pro Val Glu Leu Leu Arg Leu Asp Leu
	50 55 60
	Met Thr Pro Tyr Leu Asn Thr Ser Asn Arg Glu Val Lys Val Tyr Val
60	65 70 75 80
vv	

Cys Xaa Ile Trp Glu Asp Leu Thr Ala Ile Pro Phe Trp Val Ser Tyr 85 90 Val Pro 5 (2) INFORMATION FOR SEQ ID NO: 358: 10 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 78 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 358: Met Phe Gly Ala His Arg Xaa Trp Gln Gly Ser Val Leu Leu Phe Leu 20 Ser Phe Ala Trp Gly Asn Gly Gly Ser Val Thr Phe Ser Asp Val Pro Arg Val Met Pro Leu Ala Gly Gly Pro Xaa Xaa Gln Val Ser Ser Thr 35 40 25 Pro Arg Pro Pro Pro His Gln Val Thr Ser Ser Pro Gly Leu Glu Ser Ala His Ile Val Cys Pro Glu Arg Lys Lys Lys Lys Lys 30 (2) INFORMATION FOR SEQ ID NO: 359: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 359: Thr Leu Leu Xaa Phe Leu Xaa Leu Leu Thr Thr Glu Gly Gly Arg Glu 5 45 Așn Ile Phe Xaa Gly Arg Ile Leu Xaa Leu Gln Xaa Ser Pro Xaa 20 25 50 (2) INFORMATION FOR SEQ ID NO: 360: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 57 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 360: Met Leu Ser Phe Phe Ile Cys Leu Leu Ile Phe Val His Leu Leu Leu 10 60

Leu Ser Phe Leu Ile Ser Asp Trp Pro Pro Pro Thr Gly Ser Ala Xaa 20 His Lys Ile Leu Arg Leu Met Val Val Gln Arg Leu Ser Leu Leu Asp 5 40 Gln Arg Lys Arg Trp Ser Glu Ala Xaa 50 55 10 (2) INFORMATION FOR SEQ ID NO: 361: (i) SEQUENCE CHARACTERISTICS: 15 (A) LENGTH: 3 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 361: 20 Lys Tyr Xaa 1 25 (2) INFORMATION FOR SEQ ID NO: 362: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 32 amino acids (B) TYPE: amino acid 30 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 362: Trp Ser Ser Ala Ser Ser Ser Trp Val Thr Thr Pro Glu Arg Ile Arg 10 35 Pro Arg Met Asp Thr Leu Pro Val Lys Gly His Phe Leu Ser Met Xaa 25 40 (2) INFORMATION FOR SEQ ID NO: 363: 45 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 28 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 363: Asp Ile Phe Val Phe Leu Leu Ser Thr Arg Ala Gly Gly Leu Gly Ile 1 . 5 10 55 Asn Leu Thr Ala Xaa Asp Thr Val His Phe Leu Xaa 20 60 (2) INFORMATION FOR SEQ ID NO: 364:

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 15 amino acids
                    (B) TYPE: amino acid
 5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 364:
     Thr Leu Thr Ser Phe Leu Glu Leu Pro Leu Ala Pro Glu Pro Xaa
10
      (2) INFORMATION FOR SEQ ID NO: 365:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 34 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 365:
20
      Met His Arg Tyr Ile Thr Phe Phe Lys Cys Phe Arg Ser Val Ile Leu
                       5
                                           10
      Asp Leu Leu Phe Ile Leu Ser Pro Leu Ser Gln Gly Cys Phe Ile Leu
25
                  20
                                      25
      Phe Xaa
30
      (2) INFORMATION FOR SEQ ID NO: 366:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 66 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 366:
40
      Met Phe Gly Phe Ile Phe Leu Leu Ile Phe Cys Ile Xaa Leu Cys
                                          10
      Ser Arg Thr Leu Ser Thr Phe Ile Pro Lys Leu Val Gly Phe Leu Tyr
                   20
45
      Trp Lys Phe Ser Ile Asn Leu Ser Leu Leu Leu Thr Leu Ile Lys Lys
               35
                                   40
      Lys Lys Lys Lys Lys Thr Pro Arg Gly Gly Pro Gly Xaa Gln Ser
50
      Pro Pro
       65
55
      (2) INFORMATION FOR SEQ ID NO: 367:
             (i) SEQUENCE CHARACTERISTICS:
60
                     (A) LENGTH: 317 amino acids
```

•						YPE:										
	•		(xi)	SEQ		OPOI E DE				EQ I	D NO	: 36	7:			
5	Met 1	Pro	Gly	Leu	Gly 5	Arg	Pro	Arg	Gln	Ala 10	Arg	Trp	Thr	Leu	Met 15	Leu
10	Leu	Leu	Ser	Thr 20	Ala	Met	Tyr	Gly	Ala 25	His	Ala	Pro	Leu	Leu 30	Ala	Leu
	Cys	His	Val 35	Asp	Gly	Arg	Val	Pro 40	Phe	Arg	Pro	Ser	Ser 45	Ala	Val	Leu
15	Leu	Thr 50	Glu	Leu	Thr	Lys	Leu 55	Leu	Leu	Cys	Ala	Phe 60	Ser	Leu	Leu	Val
	Gly 65	Trp	Gln	Ala	Ттр	Pro 70	Gln	Gly	Pro	Pro	Pro 75	Trp	Arg	Gln	Ala	Ala 80
20	Pro	Phe	Ala	Leu	Ser 85	Ala	Leu	Leu	Tyr	Gly 90	Ala	Asn	Asn	Asn	Leu 95	Val
25	Ile	Tyr	Leu	Gln 100	Arg	Туг	Met	Asp	Pro 105	Ser	Thr	Tyr	Gln	Val 110	Leu	Ser
	Asn	Leu	Lys 115	Ile	Gly	Ser	Thr	Ala 120	Val	Leu	Tyr	Cys	Leu 125	Cys	Leu	Arg
30	His	Arg 130	Leu	Ser	Val	Arg	Gln 135	Gly	Leu	Ala	Leu	Leu 140	Leu	Leu	Met	Ala
	Ala 145	Gly	Ala	Cys	Tyr	Ala 150	Ala	Gly	Gly	Leu	Gln 155	Val	Pro	Gly	Asn	Thr 160
35	Leu	Pro	Ser	Pro	Pro 165	Pro	Ala	Ala	Ala	Ala 170	Ser	Pro	Met	Pro	Leu 175	His
40	Ile	Thr	Pro	Leu 180	Gly	Leu	Leu	Leu	Leu 185	Ile	Leu	Tyr	Суз	Leu 190	Ile	Ser
	Gly	Leu	Ser 195	Ser	Val	Туг	Thr	Glu 200	Leu	Leu	Met	Lys	Arg 205	Gln	Xaa	Leu
45	Pro	Leu 210	Ala	Leu	Gln	Asn	Leu 215	Phe	Leu	Tyr	Thr	Phe 220	Gly	Val	Leu	Leu
	Asn 225	Leu	Gly	Leu	His	Ala 230	Gly	Gly	Gly	Ser	Gly 235	Pro	Gly	Leu	Leu	Glu 240
50	Gly	Phe	Ser	Glý	Trp 245	Ala	Ala	Leu	Val	Val 250	Leu	Ser	Gln	Ala	Leu 255	Asn
55	Gly	Leu	Leu	Met 260	Ser	Ala	Val	Met	Lys 265	His	Gly	Ser	Ser	Ile 270	Thr	Arg
	Leu	Phe	Val 275	Val	Ser	Cys	Ser	Leu 280	Val	Val	Asn	Ala	Val 285	Leu	Ser	Ala
60	Val	Leu	Leu	Arg	Leu	Gln	Leu	Thr	Ala	Ala	Phe	Phe	Leu	Ala	Thr	Leu

•	
	Leu Ile Gly Leu Ala Met Arg Leu Tyr Tyr Gly Ser Arg 305 310 315
5	
	(2) INFORMATION FOR SEQ ID NO: 368:
10	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 368:
15	Met Gly Glu Gln Pro His Phe Ser Leu Cys Val Leu Leu Ala Ala Val 1 5 10 15
20	Arg Glu Asp Xaa Asp Pro Xaa Val Phe Pro Cys Cys Phe Leu Xaa 20 25 30
	(2) INFORMATION FOR SEQ ID NO: 369:
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 369:
50	Met Ser Phe Ile Ala Leu His Pro Leu Leu Pro Glu Ala Ala Leu Gly 1 5 10 15
35	Val Pro Gly Gln Ser Pro His Arg Pro Leu Trp Gln Thr Gln Cys Cys 20 25 30
	Val Ala Pro Pro Gln Pro Arg Ala Glu Phe Xaa 35 40
40	
	(2) INFORMATION FOR SEQ ID NO: 370:
45	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 255 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 370:
50	Met Val Thr Ala Leu Thr Leu Leu Ala Phe Pro Leu Leu Leu His 1 5 10 15
• 6 6	Ala Glu Arg Ile Ser Leu Val Phe Leu Leu Phe Leu Gln Ser Phe 20 25 30
55	Leu Leu Leu His Leu Leu Ala Ala Gly Ile Pro Val Thr Thr Pro Gly 35 40 45
60	Pro Phe Thr Val Pro Trp Gln Ala Val Ser Ala Trp Ala Leu Met Ala 50 55 60

	Thr 65	Gln	Thr	Phe	Tyr	Ser 70	Thr	Gly	His	Gln	Pro 75	Val	Phe	Pro	Ala	Ile 80
5	His	Trp	His	Ala	Ala 85	Phe	Val	Gly	Phe	Pro 90	Glu	Gly	His	Gly	Ser 95	Суз
	Thr	Trp	Leu	Pro 100	Ala	Leu	Leu	Val	Gly 105	Ala	Asn	Thr	Phe	Ala 110	Ser	His
-	Leu	Leu	Phe 115	Ala	Val	Gly	Cys	Pro 120	Leu	Leu	Leu	Leu	Trp 125	Pro	Phe	Leu
15	Cys	Glu 130	Ser	Gln	Gly	Leu	Arg 135	Lys	Arg	Gln	Gln	Pro 140	Pro	Gly	Asn	Glu
	Ala 145	Asp	Ala	Arg	Val	Arg 150	Pro	Glu	Glu	Glu	Glu 155	Glu	Pro	Leu	Met	Glu 160
20	Met	Arg	Leu	Arg	Asp 165	Ala	Pro	Gln	His	Phe 170	Tyr	Ala	Ala	Leu	Leu 175	Gln
25	Leu	Gly	Leu	Lys 180	Tyr	Leu	Phe	Ile	Leu 185	Gly	Ile	Gln	Ile	Leu 190	Ala	Суѕ
	Ala	Leu	Ala 195	Ala	Ser	Ile	Leu	Arg 200	Arg	His	Leu	Met	Val 205	Trp	Lys	Val
30	Phe	Ala 210	Pro	Lys	Phe	Ile	Phe 215	Glu	Ala	Val	Gly	Phe 220	Ile	Val	Ser	Ser
	Val 225	Gly	Leu	Leu	Leu	Gly 230	Ile	Ala	Leu	Val	Met 235	Arg	Val	Asp	Gly	Ala 240
35	Val	Ser	Ser	Trp	Phe 245	Arg	Gln	Leu	Phe	Leu 250	Ala	Gln	Gln	Arg	Xaa 255	
40	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: I	371:							
			(i)		A) L	CHA ENGT YPE:	H: 2	0 an	ino		is					
45			(xi)		D) I	OPOL	OGY:	lin	ear	EQ I	D NO	: 37	1:			
50	Met 1		Gly	Pro	Trp 5	Ģly	Glu	Glu	Ala	Leu 10	Ile	Arg	Leu	Pro	Thr 15	Pro
	Ser	Gly	Leu	Хаа 20												
55	(2)	INF	ORMA	TION	FOR	SEQ	ID :	NO:	372:							
			(i)	SEQU	ENCE	СНА	RACT	ERIS	TICS	:						
50						ENGI				acid	ls					
					ו נכו	TED:	- cuul	ة بسد	C-LU							

```
(D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 372:
       Met Ala Thr Leu Glu Xaa Asn Gln Arg Glu Val Asp Arg Glu Ile Arg
  5
                                10
       Ser Leu Leu Trp Phe Leu Leu Cys Glu Ile Val Ser Gly Trp Leu
 10
      Cys Pro Glu Gly Pro Trp Phe Ser Gln Gly Cys Gln Ile Tyr Lys Asn
               35
                                   40
      Leu Ser Ser Ser Ser Tyr Asn Leu Ser Phe Leu Leu Ser Leu Xaa
                               55
                                                   60
15
20
       (2) INFORMATION FOR SEQ ID NO: 373:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 40 amino acids
25
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 373:
      Met Ile His Ser Gly Cys Thr Ser Gln Cys Leu Glu Gly Phe Phe Leu
30
      Ile Phe Leu Leu Asp Phe Asn Pro Val Leu Ala Leu Asp Leu Ile Gly
                                     25
35
      Ile Met Arg Lys Ala Ser His Xaa
               35
40
      (2) INFORMATION FOR SEQ ID NO: 374:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 35 amino acids
                    (B) TYPE: amino acid
45
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 374:
     Met Val Phe Ser Ala Arg Val Ser Leu Tyr Thr Arg Phe Lys Val Ile
50
     Leu Leu Ser Leu Leu Ile Met Ile Leu His Val Cys Trp Val Trp Val
     Ile Leu Xaa
55
              35
```

(2) INFORMATION FOR SEQ ID NO: 375:

```
(i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 11 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
  5
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 375:
       Gly Leu Leu Tyr Ile Met Tyr Cys Asn Ile Xaa
 10
       (2) INFORMATION FOR SEQ ID NO: 376:
              (i) SEQUENCE CHARACTERISTICS:
 15
                     (A) LENGTH: 64 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 376:
 20
      Met Asn Asn Gly Leu Leu Gln Gln Pro Ser Ala Leu Met Leu Leu Pro
                                           10
      Cys Arg Pro Val Leu Thr Ser Val Ala Leu Asn Ala Asn Phe Val Ser
                                      25
 25
      Trp Lys Ser Arg Thr Lys Tyr Thr Ile Thr Pro Val Lys Met Arg Lys
      Ser Gly Gly Arg Asp His Thr Gly Gly Asn Lys Asp Arg Gly Ile Xaa
30
           50
                                55
                                                    60
35
      (2) INFORMATION FOR SEQ ID NO: 377:
              (i) SEQUENCE CHARACTERISTICS:
40
                     (A) LENGTH: 19 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 377:
45
      Met Arg Lys Gln Arg Leu Val Pro Met Tyr Leu Gly Leu Ile Tyr Ile
                        5
                                           10
      Leu Leu Xaa
50
      (2) INFORMATION FOR SEQ ID NO: 378:
55
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 5 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 378:
60
```

```
Met Arg Gln His Xaa
 5
       (2) INFORMATION FOR SEQ ID NO: 379:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 17 amino acids
10
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 379:
      Leu Leu Pro Val Leu Ala Ser Ser Val Pro Ser His Ser Ala Thr
15
                        5
                                           10
      Xaa
20
      (2) INFORMATION FOR SEQ ID NO: 380:
             (i) SEQUENCE CHARACTERISTICS:
25
                     (A) LENGTH: 84 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 380:
30
      Met Leu Pro Leu Leu Leu Phe Thr Tyr Leu Asn Ser Phe Leu His Gln
                                          10
      Arg Ile Pro Gln Ser Val Arg Ile Leu Gly Ser Leu Val Ala Ile Leu
                                       25
.35
      Leu Val Phe Leu Ile Thr Ala Ile Leu Val Lys Val Gln Leu Asp Ala
                          _ 40
      Leu Pro Phe Phe Val Ile Thr Met Ile Lys Ile Val Leu Ile Asn Ser
40
      Phe Gly Ala Ile Leu Gln Gly Ser Leu Phe Gly Leu Ala Gly Leu Leu
                          70
                                          75
45
      Pro Ala Ser Xaa
50
      (2) INFORMATION FOR SEQ ID NO: 381:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 21 amino acids
                    (B) TYPE: amino acid
55
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 381:
     Met Lys Leu Ser Leu Phe Leu Ile Leu Ser Asp Val Phe Tyr Leu Gly
                      5
                                          10
```

Ser Pro Xaa Thr Xaa 20

5

- (2) INFORMATION FOR SEQ ID NO: 382:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 29 amino acids
- 10 (B) TYPE: amino acid
 - (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 382:
- Met Gly Thr Arg Arg Lys Gly Val Ala Trp Leu Ser Leu Ala Pro Leu 15 5 10

Ile Thr Gly Leu Ala Pro Ala His Ile Thr Ala Val Xaa 20 25

20

- (2) INFORMATION FOR SEQ ID NO: 383:
- (i) SEQUENCE CHARACTERISTICS: 25
 - (A) LENGTH: 34 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 383:
- 30 Met Lys Asp Leu Leu Gln Arg Asn Pro Trp Lys Asn Ser Leu Leu Leu

Leu Gln Val Cys Gln Ala Phe Leu Val Cys Ser Leu Thr Gln Leu Ala 20 . 25

35

Val Xaa

40

45

- (2) INFORMATION FOR SEQ ID NO: 384:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 47 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 384:
- Met Ser Glu Ser His Lys Ile Trp Trp Cys Tyr Arg His Leu Ala Phe 50 · 5

Pro Leu Leu Thr Leu Ile Leu Tyr Pro Ala Thr Leu Gly Arg Ser Val

- 55 Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Xaa 35 40
- 60 (2) INFORMATION FOR SEQ ID NO: 385:

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 25 amino acids
                     (B) TYPE: amino acid
 5
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 385:
      Met Leu Asn Arg Ile Met Val Ala Ser Phe Gly Ala Val Leu Val Gln
                                 10
10
      Val Cys Arg Gly Kaa Gly Gln Gly Xaa
                   20
15
      (2) INFORMATION FOR SEQ ID NO: 386:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 68 amino acids
20
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 386:
      Met Gln Leu Leu Leu Gly Leu Ile Arg Ser Gln Pro Ser Pro Pro
25
      Pro Ser Leu Cys Leu Met Leu Cys Pro Cys Leu Pro Cys Leu Arg Tyr
                  20
                                      25
30
      Ser Pro Phe Val Pro Gln His Pro Cys Pro Leu Pro Leu Asp Leu Cys
      Leu Ala Gly Cys Ser Ser Leu Ser Val Gln Asp Lys Cys Ser Trp Pro
                                                60
35
      Tyr Pro Ile Xaa
       65
40
      (2) INFORMATION FOR SEQ ID NO: 387:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 34 amino acids
45
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 387:
     Lys Glu Phe Phe Val Phe Leu Phe Val Cys Leu Phe Trp Leu Leu Ser
50
                                         10
     Asn Thr Pro Leu Thr Phe Ile Ser Ile Ile Leu Gln Arg Lys Glu Thr
                              25
55
     Asn Xaa
60
     (2) INFORMATION FOR SEQ ID NO: 388:
```

	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 14 amino acids	
•	(B) TYPE: amino acid	
5	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 388:	
	Ser Phe Leu Met Val Leu Val Ile Leu Ala Ala Ser Pro Xaa	
	1 5 10	
10		
	(2) INFORMATION FOR THE TRANSPORT	
	(2) INFORMATION FOR SEQ ID NO: 389:	
15	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: amino acids	
	(B) TYPE: amino acid	
	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 389:	
20	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
	1 5 10	
26		
25		
	(2) INFORMATION FOR SEQ ID NO: 390:	
	III grantan and a second	
	(i) SEQUENCE CHARACTERISTICS:	
30	(A) LENGTH: 154 amino acids	
	(B) TYPE: amino acid	
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 390:	
	(mr. posomich proceditation: 250 ID MO: 330:	
	Met Thr Lys Ala Arg Leu Phe Arg Leu Trp Leu Val Leu Gly Ser Val	
35	1 5 10 15	
	Phe Met Ile Leu Leu Ile Ile Val Tyr Trp Asp Ser Ala Gly Ala Ala	
	20 25 30	•
40		
40	His Phe Tyr Leu His Thr Ser Phe Ser Arg Pro His Thr Gly Pro Pro	,
	35 40 45	
	I ou Dro Mhu Du où o	
	Leu Pro Thr Pro Gly Pro Asp Arg Asp Arg Glu Leu Thr Ala Asp Ser	
45	50 55 60	
	Asp Val Asp Yaa Dho Lou Am Yaa Bi	
	Asp Val Asp Xaa Phe Leu Asp Xaa Phe Leu Ser Ala Gly Val Lys Gln 65 70 75 80	
	70 75 80	
	Ser Asp Xaa Pro Arg Lys Glu Thr Glu Gln Pro Pro Ala Pro Gly Ser	
50	95 🛕	
	90 95	
	Met Glu Glu Ser Val Arg Xaa Tyr Asp Trp Ser Pro Arg Xaa Ala Arg	
	100 105 110 Alg Ala Alg	
	110	
5 5	Arg Thr Gln Thr Arg Ala Gly Ser Xaa Arg Xaa Gly Gly Xaa Cys Cys	
	115 120 125	
	Gly Ala Ser Ala Pro Xaa Pro Ala Trp Pro Ser Pro Pro Arg Ser Ala	
60	130 135 140	
-		

```
His Ser Thr Thr Ser Pro Thr Arg Ser Xaa
     145
                         150
5
      (2) INFORMATION FOR SEQ ID NO: 391:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 9 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 391:
     Met Val Leu Leu Gly Leu Leu Ser Xaa
15
      (2) INFORMATION FOR SEQ ID NO: 392:
20
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 61 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
25
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 392:
     Met Cys Ile His Val Phe Met Xaa Val Leu Trp Val Leu Phe Leu Leu
                  5 .
                                 10
30
     Asn Pro Leu Cys Thr Gly Leu Trp Pro Leu Xaa Asn Cys Phe Ser Val
                                      25
     Leu Arg His Ala Asp Trp Val Leu Gly Ala Asp Tyr Lys Gly Glu Glu
35
     Leu Asn Arg His Gln Gly Pro Met Lys Pro Lys Asp Xaa
40
      (2) INFORMATION FOR SEQ ID NO: 393:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 447 amino acids
45
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 393:
     Met Leu Leu Gly Leu Leu Met Ala Ala Cys Phe Thr Phe Cys Leu Ser
50.
     His Gln Asn Leu Lys Glu Phe Ala Leu Thr Asn Pro Glu Lys Ser Ser
55
     Thr Lys Glu Thr Glu Arg Lys Glu Thr Lys Ala Glu Glu Glu Leu Asp
                                 40
     Ala Glu Val Leu Glu Val Phe His Pro Thr His Glu Trp Gln Ala Leu
                              55
60
```

	Gln 65	Pro	Gly	Gln	Ala	Val 70	Pro	Ala	Gly	Ser	His 75	Val	AIG	Leu	Ast.	20 30
5	Gln	Thr	Gly	Glu	Arg 85	Glu	Ala	Lys	Leu	Gln 90	Tyr	Glu	qaA	Lys	?he 95	æş
	Asn	Asn	Leu	Lys 100	Gly	Lys	Arg	Leu	Asp 105	Ile	Asn	Thr	Asn	Thr 110	T}=	Tita
10	Ser	Gln	Asp 115	Leu	Lys	Ser	Ala	Leu 120	Ala	Lys	Phe	Lys	Glu 125	Gly	Ala	3lu
15	Met	Glu 130	Ser	Ser	Lys	Glu	Asp 135	Lys	Ala	Arg	Gln	Ala 140	Glu	Val	Lys	Arg
	Leu 145	Phe	Arg	Pro	Ile	Glu 150	Glu	Leu	Lys	Lys	Asp 155	Phe	çzA	Glu	Leu	Asn 165
20	Val	Val	Ile	Glu	Thr 165	Asp	Met	Gln	Ile	Met 170	Val	Arg	Leu	Ile	Asn 175	Lys
	Phe	Asn	Ser	Ser 180	Ser	Ser	Ser	Leu	Glu 185	Glu	Lys	Ile	Ala	Ala 190	Leu	Phe
25	Asp	Leu	Glu 195	Tyr	Тут	Val	His	Gln 200	Met	Asp	Asn	Ala	Gln 205	λsp	Leu	Leu
30	Ser	Phe 210	Gly	Gly	Leu	Gln	Val 215	Val	Ile	Asn	Gly	Leu 220	Asn	Ser	T==	31u
	Pro 225	Leu	Val	Lys	Glu	Тут 230	Ala	Ala	Phe	Val	Leu 235	Gly	Ala	λla	Phe	Ser 240
35	Ser	Asn	Pro	Lys	Val 245	Gln	Val	Glu	Ala	Ile 250	Glu	Gly	Gly	Ala	Le: 255	3ln
	Lys	Leu	Leu	Val 260	Ile	Leu	Ala	Thr	Glu 265	Gln	Pro	Leu	Thr	Ala 270	Lys	<u> -:</u> /s
40	Lys	Val	Leu 275	Phe	Ala	Leu	Cys	Ser 280	Leu	Leu	Arg	His	Phe 285	Pro	בעה	λla
45	Gln	Arg 290	Gln	Phe	Leu	Lys	Leu 295	Gly	Gly	Leu	Gln	Val 300	Leu	Arg	<u>Thr</u>	Ľеч
	Val 305	Gln	Glu	Lys	Gly	Thr 310	Glu	Val	Leu	Ala	Val 315	Arg	Val	Val	Time:	1.eu 320
50	Leu	Tyr	Asp	Leu	Val 325	Thr	Glu	Lys	Met	Phe 330	Ala	Glu	Glu	Glu	Ala 335	Glu
	Leu	Thr	Gln	Glu 340	Met	Ser	Pro	Glu	Lys 345	Leu	Gln	Gln	Tyr	Arg 350	Glm	Val
55	His	Leu	Leu 355		Gly	Leu	Trp	Glu 360	Gln	Gly	Trp	Cys	Glu 365	Ile	The	Аlа
60	His	Leu 370		Ala	Leu	Pro	Glu 375	His	Asp	Ala	Arg	Glu 380	Lys	Val	Le:	Glin

•	Thr Leu Gly Val Leu Leu Thr Thr Cys Arg Asp Arg Tyr Arg Gln As 385 390 395 40	ą 00
5	Pro Gln Leu Gly Arg Thr Leu Ala Ser Leu Glm Ala Glu Tyr Gln Va 405 410 415	al
	Leu Ala Ser Leu Glu Leu Gln Asp Gly Glu Asp Gly Gly Tyr Phe G 420 425 430	ln
10	Glu Leu Cly Ser Val Asn Ser Leu Leu Lys Glu Leu Arg Xaa 435 443 445	
15	(2) INFORMATION FOR SEQ ID NC: 394:	
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 394:	
25	Met Val Ile Ser Tyr Val Thr Phe Thr Pro Val Ser Ala Asp Cys P 1 5 10 15 Phe Asn Val Leu Val Cys Phe Kaa 20	he
30	(2) INFORMATION FOR SEQ ID NC: 395:	
35	(i) SEQUENCE CHARACTEFISTICS: (A) LENGTH: 24 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 395:	
40	Glu Leu Leu Phe Leu Leu Ile Ile Ile Leu Gly Glu Ser Leu Ser I 1 5 10 15	Asp
٠.	Val Ile Leu Leu Ile Cys Phe Kaa 20	
45	(2) THEODINAMION FOR CO. ID NO. 206.	
50	(2) INFORMATION FOR SEQ ID NC: 396: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 35 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 396:	
55	Met Phe Tyr Trp Gly Gly Leu Ser Phe Tyr Phe Leu Leu Ser Ser 1 5 10 15	Gly
60	Val Gly Phe Tyr Cys Phe Leu Phe Gly Phe Gly Met Glu Ile Trp 20 25 30	Ile

Ala Ala Xaa 35

5

- (2) INFORMATION FOR SEQ ID NO: 397:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 3 amino acids
- 10 (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 397:
 - Gly Arg Xaa
- 15
- (2) INFORMATION FOR SEQ ID NO: 398:

20

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 25 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
- 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 398:

Met Lys Leu Ser Leu Leu Ile Leu Thr Leu Met Gln Arg Tyr Phe Arg 1 5 10 15

- 30 Thr Ile Thr Asn Ser Leu Cys Lys Xaa
- 35 (2) INFORMATION FOR SEQ ID NO: 399:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 79 amino acids
 - (B) TYPE: amino acid
- 40 (D) TOPOLOGY: linear
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 399:

Met Pro Ala Val Ser Gly Pro Gly Pro Leu Phe Cys Leu Leu Leu 1 5 10 15

45

Leu Leu Asp Pro His Ser Pro Glu Thr Gly Cys Pro Pro Leu Arg Arg
20 25 30

Phe Glu Tyr Lys Leu Ser Phe Lys Gly Pro Arg Leu Ala Leu Pro Gly 50 35 40 45

Ala Gly Ile Pro Phe Trp Ser His His Gly Gly Glu Gly Gln Gly Trp 50 55 60

- 55 Gly Pro Leu Cys Pro Gly Ser Leu Lys Val Leu Glu Gly Leu Xaa 65 70 75
- 60 (2) INFORMATION FOR SEQ ID NO: 400:

PCT/US98/11422 WO 98/54963

5			(i) S	- (1 (1	A) LE 3) TY 0) TY	ENGTI (PE : OPOLA	i: 2: amir XGY:	l ami no ac line	ino a cid ear	acids		. 400	١.			
10	Met 1		(xi) Val	_						-				Gln	Ser 15	Leu
	Ile	Gln	Glu	Asp 20	Xaa											
15	(2)		ORMAT	EQUI	ENCE	CHAI	RACTI	RIST	rics							
20			(xi)	() ()	A) LI B) T D) T JENCI	YPE: OPOL	amii OGY :	no ad lind	cid ear			: 40:	1:			
25	Met 1	Ala	Ala	Leu	Thr 5	Ser	His	Leu	Gln	Asn 10	Gln	Ser	Asn	Asn	Ser 15	Asn
	Trp	Asn	Leu	Arg 20	Thr	Arg	Ser	Lys	Cys 25	Lys	Lys	Asp	Val	Phe 30	Met	Pro
30	Pro	Ser	Ser 35	Ser	Ser	Glu	Leu	Gln 40	Glu	Ser	Arg	Gly	Leu 45	Ser	Asn	Phe
35	Thr	Ser 50	Thr	His	Leu	Leu	Leu 55	Lys	Glu	Asp	Glu	Gly 60	Val	Asp	Asp	Val
	Asn 65		Arg	Lys	Val	Arg 70	Lys ·	Pro	Lys	Gly	Lys 75	Val	Thr	Ile	Leu	Lys 80
40	Gly	Ile	Pro	Ile	Ľуз 85	Lys	Thr	Lys	Lys	90 Gly	Cys	Arg	Lys	Ser	Cys 95	Ser
٠.	Gly	Phe	Val	Xaa 100	Ser	Asp	Ser	Lys	Arg 105	Glu	Ser	Val	Cys	Asn 110	Lys	Ala
45	Asp	Ala	Glu 115	Ser	Glu	Pro	Val	Ala 120	Gln	Lys	Ser	Gln	Leu 125	Asp	Arg	Thr
50	Val	Cys 130	Ile	Ser	Asp	Ala	Gly 135	Ala	Суз	Gly	Glu	Thr 140		Ser	Val	Thr
	Ser 145		Glu	Asn	Ser	Leu 150	Val	Lys	Lys	Lys	Glu 155	Arg	Ser	Leu	Ser	Ser 160
55	Gly	Ser	Asn	Phe	Cys 165	Ser	Glu	Gln	Lys	Thr 170	Ser	Gly	Ile	Ile	Asn 175	
	Phe	Cys	Ser	Ala 180	Lys	Asp	Ser	Glu	His 185	Asn	Glu	Lys	Tyr	Glu 190		Thr
60	Phe	Leu	Glu	Ser	Glu	Glu	Ile	Gly	Thr	Lys	Val	Glu	Val	Val	Glu	Arg

•			195					200					205			
5	Lys	Glu 210	His	Leu	His	Thr	Asp 215	Ile	Leu	Lys	Arg	Gly 220	Ser	Glu	Met	Asp
	Asn 225	Asn	Cys	Ser	Pro	Thr 230	Arg	Lys	Asp	Phe	Thr 235	Glu	Asp	Thr	Ile	Pro 240
10	Arg	Asn	Thr	Asp	Arg 245	Lys	Lys	Glu	Asn	Lys 250	Pro	Val	Phe	Phe	Gln 255	Gln
15	Ile															
13	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	1 0: 4	102:							
20			(i) : (xi)	() ()	A) L B) T D) T	ENGT YPE : OPOL	H: 4 ami OGY:	24 a no a lin	mino cid ear	aci		: 40	2:			
25	Met 1	Glu	Lys	Gln	Cys 5	Cys	Ser	His	Pro	Val 10	Ile	Cys	Ser	Leu	Ser 15	Thr
30	Met	Tyr	Thr	Phe 20	Leu	Leu	Gly	Ala	Ile 25	Phe	Ile	Ala	Leu	Ser 30	Ser	Ser
	Arg	Ile	Leu 35	Leu	Val	Lys	Tyr	Ser 40	Ala	Asn	Glu	Glu	Asn 45	Lys	Tyr	Asp
35	Tyr	Leu 50	Pro	Thr	Thr	Val	Asn 55	Val	Суз	Ser	Glu	Leu 60	Val	Lys	Leu	Val
10	Phe 65	Cys	Val	Leu	Val	Ser 70	Phe	Cys	Val	Ile	Lys 75	Lys	Asp	His	Gln	Ser 80
40		Asn			85					90					95	
45				100					105					110		Phe
		Val	115					120					125			
50		Ser 130					135					140				
	145					150					155					160
55		Val			165					170					175	
60	Gly	Arg	Gly	Phe 180	His	His	Asp	Ala	Phe 185	Phe	Ser	Pro	Ser	Asn 190	Ser	Cys.

	Leu	Leu	Phe 195	Arg	Asn	Glu	Cys	Pro 200	Arg	Lys	Asp	Asn	Cys 205	Thr	Ala	Lys	
5	Glu	Trp 210	Thr	Phe	Pro	Glu	Ala 215	Lys	Trp	Asn	Thr	Thr 220	Ala	Arg	Val	Phe	
	Ser 225	His	Ile	Arg	Leu	Gly 230	Met	Gly	His	Val	Leu 235	Ile	Ile	Val	Gln	Cys 240	
10	Phe	Ile	Ser	Ser	Met 245	Ala	Asn	Ile	Tyr	Asn 250	Glu	Lys	Ile	Leu	Lys 255	Glu	
15	Gly	Asn	Gln	Leu 260	Thr	Glu	Хаа	Ile	Phe 265	Ile	Gln	Asn	Ser	Lys 270	Leu	Tyr	
13	Phe	Phe	Gly 275	Ile	Leu	Phe	Asn	Gly 280		Thr	Leu	Gly	Leu 285		Arg	Ser	
20	Asn	Arg 290	Asp	Gln	Ile	Lys	Asn 295	Cys	Gly	Phe	Phe	Туг 300		His	Ser	Ala	
	Phe 305		Val	Ala	Leu	Ile 310		Val	Thr	Ala	Phe 315		Gly	Leu	Ser	Val 320	
25	Ala	Phe	lle	Leu	Lys 325		Leu	Asp	Asn	Met 330		His	Val	Leu	Met 335	Ala	
30	Gln	ı Val	. Thr	Thr 340		Ile	Ile	Thr	Thr 345		. Ser	Val	. Leu	Val 350	Phe	Asp	
30	Phe	e Arg	355		Leu	Glu	Phe	9he		Glu	Ala	Pro	365		. Leu	Leu	
35	Ser	370		: Ile	туг	: Asr	Ala 375		Lys	Pro	Glr	380		Glu	тут	Ala	
	Pro 389		g Glr	ı Glu	ı Arç	390		, Ası) Lev	Sei	Gl ₃		ı Le	ı Tr <u>ı</u>	Glu	400	
40	Se	r Se	r Gly	/ Asp	Gl ₃ 40		ı Glu	ı Lev	ı Glu	410		ı Thi	r Lys	s Pro	419	s Ser	
45	Ası	p Gl	u Sei	r Ası 420		ı Ası	Thu	r Phe	e								
	(2) IN	FORM	ATIO	N FO	R SE(םו ס	NO:	403	:							
50			(i)	SEQ	(A) (B)	E CH LENG TYPE TOPO	TH: : an	33 a ino	mino acid	aci	eb.						
55			•		_	CE D											
	Me	t Tr 1	p Gl	y Gl	n Gl	y Se 5	r Gl	n Ly	s Se		s Ph O	e Se	r As	p Le		1 Phe 5	
60	G1	y Va	l Ar	g Gl 2		u Cy	s Al	a Gl	n Pr 2		r As	p Pr	o Gl		r Pr	o His	

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567

Xaa

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(2) INFORMATION FOR SEQ ID NO: 404:

(i) SEQUENCE CHARACTERISTICS:

10

- (A) LENGTH: 80 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 404:
- Met Val Gln His Ile Gln Pro Ala Ala Leu Ser Leu Leu Ala Gln Trp 15 10

Ser Thr Leu Val Gln Glu Leu Glu Ala Ala Leu Gln Leu Ala Phe Tyr 25

20

Pro Asp Ala Val Glu Glu Trp Leu Glu Glu Asn Val His Pro Ser Leu

Gln Arg Leu Gln Xaa Leu Leu Gln Asp Leu Ser Glu Val Ser Ala Pro 25

Pro Leu Pro Pro Thr Ser Pro Gly Arg Asp Val Ala Gln Asp Pro Xaa

30

- 35 (2) INFORMATION FOR SEQ ID NO: 405:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 95 amino acids
 - (B) TYPE: amino acid
- 40
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 405:

Met Leu Asn Gln Gly Tyr Ile Arg Lys Ile Ile Leu Ile Ile Leu 10

45

Gly Ser Phe Ser Ser Pro Lys Lys Ala Ile Leu Met Gly Phe Gln Asn 25

Gln Lys Lys Ala Leu Asn Glu Glu Gln Thr Thr Gly Val Pro Met Ser 50

Ile Ser Gly Lys Leu Arg Pro Ser Arg Ser Leu Asp Phe Val Gln Pro

55 Pro Arg Phe Gln Ser Gln Gln Pro Ser Ala Val Val Asp Arg Arg Gly · 70

Phe Xaa Xaa Lys Ala Ala Arg Gly Gln Glu Phe Ser Glu Ser Xaa 90

	121	TIM ()14·24.	1014	LOIC	JUZ	10 .	₩								
5			(i) :	(A) L B) T	CHAI ENGT YPE:	H: 2 ami	57 a no a	mino cid		ds					
10			(xi)							EQ I	D NO	: 40	6:			
	Met 1	Arg	Gly	Pro	Ala 5	Gln	Ala	Lys	Leu	Leu 10	Pro	Gly	Ser	Ala	Ile 15	Gln
15	Ala	Leu	Val	Gly 20	Leu	Ala	Arg	Pro	Leu 25	Val	Leu	Ala	Leu	Leu 30	Leu	Val
	Ser	Ala	Ala 35	Leu	Ser	Ser	Val	Val 40	Ser	Arg	Thr	Asp	Ser 45	Pro	Ser	Pro
20	Thr	Val 50	Leu	Asn	Ser	His	Ile 55	Ser	Thr	Pro	Asn	Val 60	Asn	Ala	Leu	Thr
25	His 65	Glu	Asn	Gln	Thr	Lys 70	Pro	Ser	Ile	Ser	Gln 75	Ile	Ser	Thr	Thr	Leu 80
	Pro	Pro	Thr	Thr	Ser 85	Thr	Lys	Lys	Ser	Gly 90	Gly	Ala	Ser	Val	Val 95	Pro
30	His	Pro	Ser	Pro 100	Thr	Pro	Leu	Ser	Gln 105	Glu	Glu	Ala	Asp	Asn 110	Asn	Glu
	Asp	Pro	Ser 115	Ile	Glu	Glu	Glu	Asp 120	Leu	Leu	Met	Leu	Asn 125	Ser	Ser	Pro
35	Ser	Thr 130	Ala	Lys	Asp	Thr	Leu 135	Asp	Asn	Gly	Asp	Tyr 140	Gly	Glu	Pro	Asp
40	Tyr 145	Asp	Trp	Thr	Thr	Gly 150	Pro	Arg	Asp	Asp	Asp 155	Glu	Ser	Asp	Asp	Thr 160
,	Leu	Glu	Glu	Asn	Arg 165	Gly	Tyr	Met	Glu	Ile 170	Glu	Gln	Ser	Val	Lys 175	Ser
45	Phe	Lys	Met	Pro 180	Ser	Ser	Asn	Ile	Glu 185	Glu	Glu	Asp	Ser	His 190	Phe	Phe
	Phe	His	Leu 195	Ile	Ile	Phe	Ala	Phe 200	Cyż	Ile	Ala	Val	Val 205	Туг	Ile	Thr
50	Tyr	His 210	Asn	Lys	Arg	Lys	Ile 215		Leu	Leu	Val	Gln 220	Ser	Arg	Lys	Tr
55	Arg 225	Asp	Gly	Leu	Суѕ	Ser 230	Lys	Thr	Val	Glu	Тут 235	His	Arg	Leu	Asp	Glr 240
	Asn	Val	Asn	Glu	Ala 245	Met	Pro	Ser	Leu	Lys 250	Ile	Thr	Asn	Asp	Tyr 255	Ile
	Dho															

5	(2)	INFO	ORMAI	'ION	FOR	SEQ	ID N	10: 4	107:							
J			(i) :		A) L	ENGT	H: 6	ERIST 23 au no a	mino		ds					
10			(xi)	•) T	OPOLA	OGY:	lin	ear	eQ II	O 1NO:	: 407	':			
	Met 1	Phe	Met	Arg	Ile 5	Ala	Lys	Ala	Туг	Ala 10	Ala	Leu	Thr	Asp	Glu 15	Glu
15	Ser	Arg	Lys	Asn 20	Trp	Glu	Glu	Phe	Gly 25	Asn	Pro	Asp	Gly	Pro 30	Gln	Ala
20	Thr	Ser	Phe 35	Gly	Ile	Ala	Leu	Pro 40	Ala	Trp	Ile	Val	Asp 45	Gln	Lys	Asn
	Ser	Ile 50		Val	Leu	Leu	Val 55	Tyr	Gly	Leu	Ala	Phe 60	Met	Val	Ile	Leu
25	Pro 65	Val	Val	Val	Gly	Ser 70	Trp	Trp	Tyr	Arg	Ser 75	Ile	Arg	Tyr	Ser	Gly 80
	Asp	Gln	Ile	Leu	Ile 85	Arg	Thr	Thr	Gln	Ile 90	Tyr	Thr	Tyr	Phe	Val 95	Tyr
30	Lys	Thr	Arg	Asn 100	Met	Asp	Met	Lys	Arg 105	Leu	Ile	Met	Val	Leu 110	Xaa	Gly
35	Ala	Ser	Glu 115	Phe	Asp	Pro	Gln	Туг 120	Asn	Lys	Asp	Ala	Thr 125	Ser	Arg	Pro
	Thr	Asp 130		Ile	Leu	Ile	Pro 135		Leu	Ile	Arg	Glu 140	Ile	Gly	Ser	Ile
40	Asn 145	Leu	Lys	Lys	Asn	Glu 150	Pro	Pro	Leu	Thr	Cys 155	Pro	Tyr	Ser	Leu	Lys 160
	Ala	Arg	Val	Leu	Leu 165	Leu	Ser	His	Leu	Ala 170	-	Met	Lys	Ile	Pro 175	Glu
45	Thr	Leu	Glu	Glu 180	Asp	Gln	Gln	Phe	Met 185		Lys	Гуз	Суз	Pro 190	Ala	Leu
50	Leu	Gln	Glu 195	Met	Val	Asn	Val	Ile 200	Cys	Gln	Leu	Ile	Val 205	Met	Ala	Arg
	Asn	Arg 210		Glu	Arg	Glu	Phe 215	-	Ala	Pro	Thr	Leu 220	Ala	Ser	Leu	Glu
55	Asn 225	_	Met	Lys	Leu	Ser 230		Met	Ala	Val	Gln 235	_	Leu	Gln	Gln	Phe 240
	Lys	Ser	Pro	Leu	Leu 245		Leu	Pro	His	Ile 250		Glu	Asp	Asn	Leu 255	Arg
60	Ara	Val	Ser	Asn	His	Lvs	Lvs	Tvr	Lvs	Tle	Lvs	Thr	Ile	Gln	Asp	Leu

				260					265					270		
~	Val	Ser	Leu 275	Lys	Glu	Ser	Asp	Arg 280	His	Thr	Leu		His 285	Phe	Leu	Glu
5	Asp	Glu 290	Lys	Tyr	Glu	Glu	Val 295	Met	Ala	Val	Leu	Gly 300	Ser	Phe	Pro	Tyr
0	Val 305	Thr	Met	Asp	Ile	Lys 310	Ser	Gln	Val	Leu	Asp 315	Asp	Glu	Asp	Ser	Asn 320
	Asn	Ile	Thr	Val	Gly 325	Ser	Leu	Val	Thr	Val 330	Leu	Val	Lys	Leu	Thr 335	Arg
15	Gĺn	Thr	Met	Ala 340	Glu	Val	Phe	Glu	Lys 345	Glu	Gln	Ser	Ile	Cys 350	Ala	Ala
20	Glu	Glu	Gln 355	Pro	Ala	Glu	Asp	Gly 360	Gln	Gly	Glu	Thr	Asn 365	Lys	Asn	Arg
LU	Thr	Lys 370	Gly	Gly	Trp	Gln	Gln 375		Ser	Lys	Gly	Pro 380	Lys	Lys	Thr	Ala
25	Lys 385		Lys	Lys	Lys	Lys 390	Pro	Leu	Lys	Lys	Lys 395	Pro	Thr	Pro	Val	Leu 400
	Leu	Pro	Gln	Ser	Lys 405		Gln	Lys	Gln	Lys 410		Ala	Asn	Gly	Val 415	Val
30	Gly	Asn	Glu	Ala 420		Val	Lys	Glu	Asp 425		Glu	Glu	Val	Ser 430	Asp	Lys
35	Gly	Ser	435		Glu	Glu	Glu	440		Asn	Arg	Asp	Ser 445	Gln	Ser	Glu
	Lys	Asr 450		Gly	Sex	Asp	Arg 455		Ser	Asp	Arg	Glu 460	Gln	Asp	Glu	Lys
40	Gln 465		ı Lys	Asp	Asp	Glu 470		a Glu	Trp	Glr	475		Gln	Gln	Ser	Ile 480
	Glr	Arg	, Lys	Glu	Arg 485		Leu	ı Leu	Glu	490		Ser	Lys	Ile	Thr 495	
45	Pro	Va!	l Tyr	Ser 500		туг	Phe	e Pro	505		lys	Gln	Glu	Trp 510		Trp
50	Leu	тул	r Ile 519		Ası	Arg	, Lys	520		Thi	: Leu	Ile	Ser 525		Pro	Туг
50	His	53		s Thr	Leu	ı Lys	539		Glu	ı Glı	ı Val	Glu 540		Lys	Phe	Pro
55	Ala 549		o Gly	/ Lys	Pro	Gl _y 550		n Tyr	r Glr	ту:	555		. Phe	: Leu	Arg	Ser 560
	Ası) Se	r Ty	r Met	Gl ₃ 569		ı Ası	p Glı	n Ile	E Ly:		Lev	Glu	ı Val	. Xaa 579	a Lys
60	Pho	e Me	t Ar	g Le	ı Ly:	s Pro	va	l Pro	o Glu	ı As	n His	s Pro	Glr	ı Trş	Ası	o Thr

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•		5	580				585					590		
5	Ala Ile	Glu (595	Gly A	sp Gl	ı Asp	Gln 600	Glu	Asp	Ser	Glu	Gly 605	Phe	Glu	Asp
J	Ser Phe 610		Gly G	ly Ar	Gly 615	Arg	Glu	Glu	Gly	Arg 620	Trp	Trp	Thr	
10	(2) INF	ORMAT:	ION F	OR SE	Q IO	NO: 4	108:							
15			(A) (B) (D)	ICE CH LENC TYPE TOPC	TH: 1 : ami LOGY:	.90 a no a lin	mino cid ear	aci		: 40	8:			
20	Met Lys 1	Ala	Ser G	ln Cy 5	s Cys	Cys	Cys	Leu 10	Ser	His	Leu	Leu	Ala 15	Ser
	Val Leu	Leu :	Leu L 20	eu Le	u Leu	Pro	Glu 25	Leu	Ser	Gly	Xaa	Leu 30	Xaa	Val
25	Leu Leu	Gln . 35	Ala A	la Gl	u Ala	Ala 40	Pro	Gly	Leu	Gly	Pro 45	Pro	Asp	Pro
30	Arg Pro	-	Thr L	æu Pr	o Pro 55		Pro	Pro	Gly	Pro 60	Thr	Pro	Ala	Gln
	Gln Pro 65	Gly .	Arg G	-	u Ala O	Glu	Ala	Ala	Gly 75	Pro	Arg	Gly	Ser	Glu 80
35	Gly Gly	Asn (Gly S	Ser As 85	n Pro	Val	Ala	Gly 90	Leu	Glu	Thr	Asp	Asp 95	His
	Gly Gly	_	Ala 0	Gly Gl	u Gly	Ser	Val 105	Gly	Gly	Gly	Leu	Ala 110	Val	Ser
40	Pro Asr	Pro 115	Gly #	Asp Ly	s Pro	Met 120	Thr	Gln	Arg	Ala	Leu 125	Thr	Val	Leu
45	Met Val		Ser (Gly Al	a Val		Val	Tyr	Phe	Val 140		Arg	Thr	Val
	Arg Met	Arg	Arg #	Arg As		Lys	Thr	Arg	Arg 155		Gly	Val	Leu	Asp 160
50	Thr Asr	lle		Asn Me 165	et Glu	Leu	Thr	Pro 170	Leu	Glu	Gln	Asp	Asp 175	
	Asp Asp) Asp	Asn 1 180	Thr Le	u Phe	Asp	Ala 185		His	Pro	Arg	Arg 190		
55							•							
	(2) IN	TORMAT	I NOI	FOR SI	Q ID	NO:	409 :							
60		(i) S	-	NCE C					.ds					

•						TPE:										
			(xi)	SEÇ	JEK	EDE	SCPI	2770	T: 5	35 =	2 110	: 40	9:			
5	Met 1	Ser	Pro	Ser	Gly S	Arg	Le:1	Ciz	Leu	Leu 10	Thr	Ile	Val	Gly	Leu 15	Ile
10	Leu	Pro	Thr	Arg 20	Gly	GLn	T <u></u> -	Leu	Lys 25	Asp	Thr	Thr	Ser	Ser 30	Ser	Ser
	Ala	ysb	Ser 35	The	Ile	Met	Asp	Ile 40	Glm	Val	320	Thr	Arg 45	Ala	Pro	Asp
15	Ala	Val 50	Ξyr	Thr	Glu	Leu	Gln 55	₽≃o	Thr	Ser	320	Thr 60	Pro	Thr	طتق	Pro
	Ala 65	qεA	Glu	Thr	Pri	31.n 70	Pro	Gln		Gln	Thr 75	Gln	Gla	Leu	Glu	Gly 80
20	Thr	ÇZA	Gly	Sto	Lei 85	Val	T:	λsp	Pro	Glu 90	Thr	His	L:/s	Ser	Thr 95	Lys
25	Ala	Ala	His	2rs 100	Thr	¥zb	Asp	Thr	135	Thr	Leu	Ser	Glu	Arg 110	5.co	Ser
	Pro	Ser	Thr 115	ನಿತರಿ	Va_	GLA	Thr	Asp 120	3.550	Gln	The	Leu	Lys 125	Pro	Ser	Gly
30		130	Glu				135					140				
25	145		Leu			150					155					160
35			Ser	Gly	Lys 165	C;·s	yzā	Gln	Leu	Ser 170	Arg	Leu	Суs	Arg	Asn 175	His
40	Cys	Arg	Xaa													
٠.	121	TVE	CRMA	~ ~ ~»;	=7=	550	TP	NT -	4-1.							
45	(2)			SEQU (ENCE (A) [. CEA .2:161 .7:73:	RACT TH:]	ERIS .4 an	TICS ino cid		is					
50			(xi) Lys	SEQ	UEV.		SCFI	PTIC	ev: S					. Xaa		
55	1				5			-		10						
	(2)	INF	ORMA			. 53Q : C3P				i:						
						EK.					.ds					
60					(3)		: 27	ه دها	acid							

	(D) TOPOLOGY: linear	
(xi)	SEQUENCE DESCRIPTION: SEQ ID NO:	411:

Met Leu Ala Gly Lys Leu Ile Pro Val His Gln Val Arg Gly Leu Lys 5 Glu Lys Ile Val Arg Ser Phe Glu Val Ser Pro Asp Gly Ser Phe Leu 25 10 Leu Ile Asn Gly Ile Ala Gly Tyr Leu His Leu Leu Ala Met Lys Thr 40 Lys Glu Leu Ile Gly Ser Met Lys Ile Asn Gly Arg Val Ala Ala Ser 55 15 Thr Phe Ser Ser Asp Ser Lys Lys Val Tyr Ala Ser Ser Gly Asp Gly Glu Val Tyr Val Trp Asp Val Asn Ser Arg Lys Cys Leu Asn Arg Phe 20 Val Asp Glu Gly Ser Leu Tyr Gly Leu Ser Ile Ala Thr Ser Arg Asn 105 25 Gly Gln Tyr Val Ala Cys Gly Ser Asn Cys Gly Val Val Asn Ile Tyr Asn Gln Asp Ser Cys Leu Gln Glu Thr Asn Pro Lys Pro Ile Lys Ala 135 30 Ile Met Asn Leu Val Thr Gly Val Thr Ser Leu Thr Phe Asn Pro Thr 150 155 Thr Glu Ile Leu Ala Ile Ala Ser Glu Lys Met Lys Glu Ala Val Arg 35 170 Leu Val His Leu Pro Ser Cys Thr Val Phe Ser Asn Phe Pro Val Ile 40 Lys Asn Lys Asn Ile Ser His Val His Thr Met Asp Phe Ser Pro Arg 195 200 Ser Gly Tyr Phe Ala Leu Gly Asn Glu Lys Gly Lys Ala Leu Met Tyr 45 Arg Leu His His Tyr Ser Asp Phe (2) INFORMATION FOR SEQ ID NO: 412: (i) SEQUENCE CHARACTERISTICS:

50

55

(A) LENGTH: 54 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 412:

Ile Leu Leu Cys Ser Trp Pro Thr Gly Leu Val Gly Gly Arg Asp Pro 60 5 10

574

Gly Ser Ser Arg Gly Ser Ser Ala Ser Leu Thr Pro Ser Pro Gly Arg Gln Pro Cys Ser Arg Arg Gly Tyr Ser Val Gly Arg Arg Ser Ser Pro Pro Asp Gly Ser Xaa 50 10 (2) INFORMATION FOR SEQ ID NO: 413: 15 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 33 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 413: 20 Met Ser Leu Gln Ser Asn Ala Trp Ser Lys Xaa Leu Phe Ile Val Phe 10 Leu Phe Leu Arg Val Leu Phe Lys Thr Gly Val Ser Ser Glu Glu Ser 25 20 25 30 (2) INFORMATION FOR SEQ ID NO: 414: (i) SEQUENCE CHARACTERISTICS: 35 (A) LENGTH: 219 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 414: 40 Met Ala Val Val Leu Leu Ala Asn Leu Ala Gln Gly Asp Ser Leu Ala Ala Arg Ala Ile Ala Val Gln Lys Gly Ser Ile Gly Asn Leu Leu Gly 45 Phe Leu Glu Asp Ser Leu Ala Ala Thr Gln Phe Gln Gln Ser Gln Ala 35 40 Ser Leu Leu His Met Gln Asn Pro Pro Phe Glu Pro Xaa Ser Val Asp 50 55 Met Met Arg Arg Ala Ala Arg Ala Leu Leu Ala Leu Ala Lys Val Asp 55 Glu Asn His Ser Glu Phe Thr Leu Tyr Glu Ser Arg Leu Leu Asp Ile 90 Ser Val Ser Pro Leu Met Asn Ser Xaa Val Ser Gln Val Ile Cys Asp 100 105 60

	Val	Leu	Phe 115	Leu	Xaa	Trp	Pro	Val 120	Met	Thr	Ala	Val	Gly 125	His	Leu	Pro
5	Pro	Pro 130	Cys	Val	Cys	Ala	Суs 135	Val	Glu	Asn	Leu	Glu 140	Thr	Asp	Cys	Cys
	Pro 145	Leu	Phe	Met	Gln	Asn 150	His	Leu	Arg	Ile	Gln 155	Phe	Thr	Leu	Суѕ	Cys 160
10	Pro	Ala	Ser	Pro	Leu 165	Gly	Lys	Ser	Leu	Ser 170	Cys	Phe	Ser	Leu	Leu 175	Leu
15	Pro	Pro	Pro	Leu 180	Pro	Pro	Ser	Pro	His 185	Ala	Phe	Leu	Phe	Leu 190	Val	Leu
	Thr	Leu	Leu 195	Pro	Ser	Gly	Pro	Tyr 200	Pro	Thr	Leu	Phe	Glu 205	Lys	Thr	Lys
20	Leu	Cys 210	Leu	His	Arg	Arg	Leu 215	Phe	Leu	Phe	Xaa					
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	NO: 4	115:							
25			(i)	SEQUI		CHA ENGT					s					
30			(xi)	-	D) T	YPE: OPOL E DE:	OGY:	lin	ear	EQ I	D NO	: 41	5:			
	Met 1	Leu	Pro	Asp	Glu 5	Ser	Phe	Gly	Leu	Leu 10	Leu	Ser	Ile	Pro	Ser 15	Leu
35	Thr	Pro	Ser	Ala 20	Ala	Ala	Pro	Ser	Phe 25	Cys	Val	His	Leu	Met 30	Gln	Ala
40	Ser	Arg	Ser 35	Ser	Lys	Arg	Ala	Ser 40	His	Val	Pro	Val	His 45	Leu	Leu	Trp
	Gly	Asp 50	Xaa													
45	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 4	116:							
				SEQU	ENCE		RACT	ERIS	TICS		s					
50			(xi)		D) T	YPE: OPOL E DE	OGY:	lin	ear	EQ I	D NO	: 41	6:			
55	Met 1	Arg	Pro	Gly	Ser 5		Ser	Phe	Ile	Ala 10	Phe	Leu	Ala	Thr	Glu 15	
		Ser	Cys	Phe 20			Arg	Pro	Asp 25		Хаа	Thr	Gly	Met 30	Trp	
60	_	_,	Leu		_											

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40 45 35 Arg Xaa 50 5 (2) INFORMATION FOR SEQ ID NO: 417: 10 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 70 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 417: 15 Asp Arg Pro Cys Pro Ser Ser Leu Trp Lys Val Phe Pro Leu Leu Leu Leu Leu Met Arg Leu Phe Pro Leu Pro Val Pro Gly Asn Gln Arg Ala 20 25 Xaa Leu Pro His Pro Phe Xaa Ala Pro Arg Leu Pro Cys Leu Leu Cys 40 25 Leu Cys Thr Gln Gln Phe Xaa Val Cys Ser His Tyr Leu Pro Ala Gly 55 Tyr Arg Val Asn Ser Xaa 30 (2) INFORMATION FOR SEQ ID NO: 418: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 40 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 418: 40 Met His Glu Lys Ala Trp Asn Leu Ile Leu Leu Trp Trp Leu Ser Leu 5 Asp Leu Leu Gly Val Ala Lys Thr Ala Met Trp Ala Gln Trp Cys Gly 45 25 Leu Asn Asp His Lys Gly Lys Xaa 35 50 (2) INFORMATION FOR SEQ ID NO: 419: (i) SEQUENCE CHARACTERISTICS: 55 (A) LENGTH: 22 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 419: 60

Met Ala Phe Val Leu Leu Xaa Cys Phe Val Xaa Leu Gln Ser Ser Xaa

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```
1
                                          20
                                                              15
     Gly Arg Ala Val Gln Xaa
                  20
 5
      (2) INFORMATION FOR SEQ ID 10: 420:
10
             (i) SEQUENCE CHARACTERISTICS:
                   (A) LENGTH: 33 amin: acids
                    (B) TYPE: amino acii
                    (D) TOPOLOGY: limear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 425:
15
     Met Phe Ser Leu Leu Trp Leu Val Cys Val Pro Ser Asn Ser Ser Val
                     5
                                         10
     Ala Asn Val Thr Ala Ser Arg Gly Gly Val Fhe Lys Arg Ser Leu Gly
20
                                    25
     His Glu Gly Phe Ser Xaa
             35
25
      (2) INFORMATION FOR SEQ ID NO: 421:
             (i) SEQUENCE CHARACTERISTICS:
30
                    (A) LENGTH: 35 amino asids
                    (B) TYPE: amino acii
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 421:
35
     Lys Trp Leu Leu Phe Ile Phe Leu Lei Cys Leu Gln Leu Val Asn Ala
     Leu Leu Ser Leu Phe Gln Glu Arg Phe Val His Cys Pro Ala Arg Phe
40
     Val Ser Xaa
              35
45
      (2) INFORMATION FOR SEQ ID 10: 422:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 32 amino acids
50
                    (B) TYPE: amino acii
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 422:
     Met Leu Leu Phe Leu Ser Ila Thr Asn Ser Leu Ser Phe Ila Ser Val
55
     Asp Lys Pro Phe Gly Gln Ser Glu Asp Val Trs Pro Val Ile Ser Xaa
                           25
60
```

5	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	423 :							
10			•	1	(A) 1 (B) 1 (D) 1	E CHA LENGI TYPE: TOPOL LE DE	M: 1 ami OGY:	27 a no a lir	mino acid aear	aci): 4 2	:3:			
15	Met 1	Glu	Phe	e Leu	Phe 5	Asn	Lys	Thr	Gly	Trp 10		Phe	Ala	Ala	Leu 15	
	Phe	Val	Leu	Ala 20	Met	Thr	Ser	Gly	Gln 25	Met	Trp	Asn	His	Ile 30	Arg	Gly
20	Pro	Pro	Тут 35		His	Lys	Asn	Pro 40		Thr	Gly	His	Val 45		Tyr	Ile
	His	Gly 50	Ser	Ser	Gln	Ala	Gln 55	Phe	Val	Ala	Glu	Thr 60	His	Ile	Val	Leu
25	Leu 65	Phe	Asn	Gly	Gly	Val 70	Thr	Leu	Gly	Met	Val 75	Leu	Leu	Cys	Glu	Ala 80
30	. Ala	Thr	Ser	Asp	Met 85	Asp	Ile	Gly	Lys	Arg 90	Lys	Ile	Met	Cys	Val 95	Ala
	Gly	Ile	Gly	Leu 100	Val	Val	Leu	Phe	Phe 105	Ser	Trp	Met	Leu	Ser 110	Ile	Phe
35	Arg	Ser	Lys 115		His	Gly	Tyr	Pro 120	Tyr	Ser	Phe	Leu	Met 125	Ser	Xaa	
40	(2)					SEQ				<u>.</u>						
45				() ()	A) L B) T D) T	ENGT YPE: OPOLA E DES	H: 6 ami OGY:	9 am no a lin	ino a cid ear	acid		: 42	4:			
	Met 1	Thr	Trp	His	Ser 5	Arg	Glu	Ser	Phe	Xaa 10	Leu	Leu	Arg	Val	Val 15	Ala
50	Pro	Ser	Gln	Ala 20	Pro	Gly	Met	Gln	Val 25	Ser	Pro	Ser	Gln	Arg 30	Ala	Trp
55	Arg	Arg	Pro 35	Leu	His	Arg	Cys	His 40	Val	Ala	Ala	Pro	Arg 45	Pro	His	His
	Phe	Ala 50	Phe	Phe	Arg	Asn	Pro 55	Phe	Ser	Trp	Ser	Phe 60	Ile	Lys	Leu	Leu
60	Tyr 65	Arg	Tyr	Leu	Xaa											

5	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	425 :							
•			(i)	((A) I (B) I	ENGI TYPE :	TH: 9	2 am	ino cid		ls					
10			(xi)	SEQ	(D) I					EQ I	D NO	: 42	5:			
	Met 1	Gly	Leu	Lys	Leu 5		Gly	Arg	Тут	Ile 10		Leu	Ile	Leu	Ala 15	Val
15	Gln	Ile	Ala	Тут 20	Leu	Val	Gln	Ala	Val 25	Arg	Ala	Ala	Gly	Lys 30	Cys	Asp
20	Ala	Val	Phe 35	Lys	Gly	Phe	Ser	Asp 40	Cys	Leu	Leu	Lys	Leu 45	Gly	Asp	Thr
	Ттр	Pro 50	Thr	Thr	Arg	Ser	Leu 55	Gly	Arg	Gln	Asp	Glu 60	His	Gln	Asp	Arg
25	Val 65	His	Ile	Leu	Gly	Gly 70	Phe	Pro	Gln	Leu	His 75	Gly	His	Ser	Pro	Туг 80
	Gly	Leu	Pro	Gly	Arg 85	Gly	Glu	Arg	Tyr	Val 90	Gly	Xaa				
30	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: 4	126:					-•		
35				(A) L B) T D) T	ENGT YPE: OPOL	H: 3 ami OGY:	80 a no a lin	mino cid ear	aci		: 42	6:			
40	Met 1	Ala	Arg	Arg	Ser 5	Ala	Phe	Pro	Ala	Ala 10	Ala	Leu	Trp	Leu	Trp 15	Ser
45	Ile	Leu	Leu	Cys 20	Leu	Leu	Ala	Leu	Arg 25	Ala	Glu	Ala	Gly	Pro 30	Pro	Gln
	Glu	Glu	Ser 35	Leu	Tyr	Leu	Trp	Ile 40	Asp	Ala	His	Gln	Ala 45	Arg	Val	Leu
50	Ile	Gly 50	Phe	Glu	Glu	Asp	Ile 55	Leu	Ile	Val	Ser	Glu 60	Gly	Lys	Met	Ala
	Pro 65	Phe	Thr	His	Asp	Phe 70	Arg	Lys	Ala	Gln	Gln 75	Arg	Met	Pro	Ala	Ile 80
55	Pro	Val	Asn	Ile	His 85	Ser	Met	Asn	Phe	Thr 90	Trp	Gln	Ala	Ala	Gly 95	Gln
50	Ala	Glu	Tyr	Phe 100	Tyr	Glu	Phe	Leu	Ser 105	Leu	Arg	Ser	Leu	Asp 110	Lys	Gly.

	Ile	Met	115	Asp	Pro	Thr	Val	120	Val	PTO	Leu	Leu	125	ınr	vai	PTO
5	His	Lys 130	Ala	Ser	Val	Val	Gln 135	Val	Gly	Phe	Pro	Cys 140	Leu	Gly	Lys	Gln
	Asp 145	Gly	Val	Ala	Ala	Phe 150	Glu	Val	Asp	Val	Ile 155	Val	Met	Asn	Ser	Glu 160
10	Gly	Asn	Thr	Ile	Leu 165	Gln	Thr	Pro	Gln	Asn 170	Ala	Ile	Phe	Phe	Lys 175	Thr
15	Cys	Gln	Gln	Ala 180	Glu	Суз	Pro	Gly	Gly 185	Cys	Arg	Asn	Gly	Gly 190	Phe	Cys
	Asn	Glu	Arg 195	Arg	Ile	Cys	Glu	Cys 200	Pro	Asp	Gly	Phe	His 205	Gly	Pro	His
20	Cys	Glu 210		Ala	Leu	Суѕ	Thr 215	Pro	Arg	Cys	Met	Asn 220	Gly	Gly	Leu	Cys
	Val 225	Thr	Pro	Gly	Phe	Cys 230	Ile	Cys	Pro	Pro	Gly 235	Phe	Tyr	Gly	Val	Asn 240
25	Cys	Asp	Lys	Ala	Asn 245	Cys	Ser	Thr	Thr	Суs 250		Asn	Gly	Gly	Thr 255	
30	Phe	Tyr	Pro	Gly 260		Cys	Ile	Xaa	Pro 265		Gly	Leu	Glu	Gly 270	Glu	Gln
	Cys	Glu	11e 275		Lys	Cys	Pro	Gln 280		Cys	Arg	Asn	Gly 285		Lys	Cys
35	Ile	Gly 290	_	Ser	Lys	Cys	Lys 295		Ser	Lys	Gly	Tyr 300		Gly	Asp	Leu
	Cys 305		Lys	Pro	Val	Cys 310	•	Pro	Gly	Cys	Gly 315		His	Gly	Thr	320
40	His	Glu	Pro	Asn	1 Lys 325		Gln	Cys	Gln	330		Trp	His	Gly	335	His
45				340)				345	,				350		Ala
	Gly	' Ala	355		Arg	Glr.	His	360		Ser	Leu	Lys	365		Glu	Glu
50	Arg	370) J Asp	Pro	Pro	Glu	375		Туг	: Ile	e Trp	380				
	(2)	INE	FORM	TION	ı FOF	R SEÇ) ID	NO:	427 :	;						
55			(i)	SEQU		E CHI					ds					
60			1-0	٠	(D)	TYPE TOPO	LOGY	: li	near		TD	∩ • 4°) 7 .			

Met Thr Ser Asn Leu Leu Leu Leu Thr Leu Leu Lys Asp Thr Leu

```
10
     Xaa Leu Ala Lys Xaa Asn Xaa Xaa
                  20
     (2) INFORMATION FOR SEQ ID NO: 428:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 47 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
15
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 423:
     Met Arg His His Thr Gln Leu Asn Phe Ile Phe Leu Val Glu Met Val
20
      Phe Leu His Val Gly Gln Ala Gly Leu Lys Leu Pro Thr Ser Gly Asp
                                     25
      Xaa Ala Cys Phe Gly Leu Pro Lys Val Leu Gly Leu Gln Ala Xaa
25
                                  40
      (2) INFORMATION FOR SEQ ID NO: 429:
30
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 5 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 429:
35
      Met Cys Ser Asp Xaa
40
       (2) INFORMATION FOR SEQ ID NO: 430:
              (i) SEQUENCE CHARACTERISTICS:
 45
                     (A) LENGTH: 144 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 430:
       Leu Leu Ser Ile Leu Leu Cys Leu Leu Ala Ser Gly Leu Val Val Phe
 50
                                      10
                       5
       Phe Leu Phe Pro His Ser Val Leu Val Asp Asp Asp Gly Ile Lys Val
                                        25
 55
       Val Lys Val Thr Phe Asn Lys Gln Asp Ser Leu Val Ile Leu Thr Ile
                                    40
       Met Ala Thr Leu Lys Ile Arg Asn Ser Asn Phe Tyr Thr Val Ala Val
 60
                               55
```

	Thr :	Ser	Leu	Ser	Ser	Gln 70	Ile	Gln	Туг	Met	Asn 75	Thr	Val	Val	Asn	Phe 80
5	Thr	Gly	Lys	Ala	Glu 85	Met	Gly	Gly	Pro	Phe 90	Ser	Tyr	Val	Tyr	Phe 95	Phe
Λ	Cys	Thr	Val	Pro 100	Glu	Ile	Leu	Val	His 105	Asn	Ile	Val	Ile	Phe 110	Met	Arg
10	Thr	Ser	Val 115		Ile	Ser	Tyr	Ile 120		Leu	Met	Thr	Gln 125	Ser	Ser	Leu
15	Glu	Thr 130		His	Tyr	Val	Asp 135		Gly	Gly	Asn	Ser 140	Thr	Ala	Ile	Xaa
20																
20	(2)	INF	ORMA	TION	FOF	SEC	Q ID	NO:	431:							
25					(A) : (B) : (D) :	LENG TYPE TOPO	TH: : am LOGY	reris 37 a ino : li :Pri	mino acid near	aci		O: 4:	31:			
30	Met 1		e Phe	e Phe		а Ту т 5	r Va	l Ty:	r Sei	c Val		ı Cy:	s Gl	y Le	Let 1	u Val
25	Tyr	Pro	Se:	r Lei		o Se	r Hi	s Se	r Va:		r Le	u Va	l Th	r Se	r Le	u Val
35	Ala	Se	r Al	a Le	u Xa	a										
40	(2)	IN	FORM	ATIO	n fo	R SE	Z IE	NO:	432	:						
			(i)	SEC				TERI								
45					(B) (D)	TYP	E: ai	37 a mino Y: 1:	ació inea:	1 r		· ·				
								RIPT						_		
50		t Al 1	a Se	er Il	.e As	sn Al	la Va	al Ty	r Il		.s Vā .0	il Ph	ie Le	eu Gi	ly Va	al Cys 15
	Va.	1 G1	ln Al		er Al	la A	la C	ys Pi		np C} 25	rs Se	er Gl	in Cy	/s Ai	rg Xa 30	aa Gly
55	Se:	r Va		ro Se 35	er Xa	aa										
60	(2) II	NFOR	MATI	ON F	OR S	EQ I	D NO	: 43	3:						

5	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 192 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 433:
10	Met Met Ala Ala Met Val Leu Thr Ser Leu Ser Cys Ser Pro Val Val 1 5 10 15
10	Gln Ser Pro Pro Gly Thr Glu Ala Asn Phe Ser Ala Ser Arg Ala Ala 20 25 30
15	Cys Asp Pro Trp Lys Glu Ser Gly Asp Ile Ser Asp Ser Gly Xaa Ser 35 40 45
	Thr Thr Ser Gly His Trp Ser Gly Ser Ser Gly Val Ser Thr Pro Ser 50 55 60
20	Pro Pro His Pro Gln Ala Ser Pro Lys Tyr Leu Gly Asp Ala Phe Gly 65 70 75 80
25	Ser Pro Gln Thr Asp His Gly Phe Glu Thr Asp Pro Asp Pro Phe Leu 85 90 95
23	Leu Asp Glu Pro Ala Pro Arg Lys Arg Lys Asn Ser Val Lys Val Met 100 105 110
30	Tyr Lys Cys Leu Trp Pro Asn Cys Gly Lys Val Leu Arg Ser Ile Val 115 120 125
	Gly Ile Lys Arg His Val Lys Ala Leu His Leu Gly Asp Thr Val Asp 130 135 140
35	Ser Asp Gln Phe Lys Arg Glu Glu Asp Phe Tyr Tyr Thr Glu Val Gln 145 150 155 160
40	Leu Lys Glu Glu Ser Ala Ala Ala Ala Ala Ala Ala Ala Ala Asp Pro 165 170 175
	Gln Ser Leu Gly Leu Pro Pro Pro Ser Gln Leu Pro Pro Pro Ala Xaa 180 185 190
45	
	(2) INFORMATION FOR SEQ ID NO: 434:
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids
55	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 434:
-	Met Ser Thr Asn Tyr Leu Thr Asp Val Cys Ser Leu Phe Ser Tyr Leu 1 5 10 15
60	Asn Tyr Leu Tyr Phe His His His Leu Pro Val Pro Asn Thr Xaa

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	20	25	30
5	(2) INFORMATION FOR SEQ 1		
10	(B) TYPE: (D) TOPOLO (xi) SEQUENCE DES	a: 101 amino ac: amino acid XGY: linear CRIPTION: SEQ	ID NO: 435:
15	1 5	10	u Tyr Leu Ala Leu Ser Arg 15 s Arg Ile Asn Phe Phe Val
20	Ala Thr Ser Tyr Phe Ser	Val Tyr Val Ar 40	g Gly Xaa Pro Xaa Val Pro 45
	Ala Asp Thr Pro Leu Gly 50	Pro Leu Leu Se 55	er Leu Trp Leu His His Asn 60
25	Ala Phe Phe Ser Ile Leu 65 70	Pro Lys Phe Pr	o Glu Asn Xaa Xaa Phe Leu 75 80
30	Ile Leu Lys Lys Leu Val 85	Val Glu Met Gl	ly Trp Asp Leu Phe Ile Ser 90 95
	Pro Glu Asn Lys Xaa 100		
35	(2) INFORMATION FOR SEQ	ID NO: 436:	
40	(B) TYPE	TH: 37 amino ac : amino acid LOGY: linear	
45	1 5		eu Val Phe Met Lys Val Ser 10 15
	Leu Asn Thr Thr Trp Pro	o Ala Pro Arg F 25	Pro Ala Thr Leu Arg Thr Ala 30
50	Asn Lys Ser Lys Xaa 35		
55	(2) INFORMATION FOR SE	Q ID NO: 437:	
		HARACTERISTICS: GTH: 42 amino a E: amino acid	cids
60		E: amino acid OLOGY: linear	

```
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 437:
     Phe Ser Thr Ile Arg Ser Gly Leu Thr Asp Arg Ser Val Asn Phe Leu
                                          10
5
     Phe Leu Phe Leu Asp Val Pro Asp Cys Arg Leu Val Asn Ile Glu Leu
     Met Ala Asn Ser Thr Val Thr His Ala Xaa
10
              35
      (2) INFORMATION FOR SEQ ID NO: 438:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 1 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 438:
20
      Leu
        1
25
       (2) INFORMATION FOR SEQ ID NO: 439:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 25 amino acids
30
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 439:
       Met Pro Trp Arg Arg Ala Gly Leu Met Met Leu Pro Ile Ile Thr Gly
 35
                         5
         1
       Cys Cys Pro Cys Ser Ala Ser Ile Xaa
                    20
 40
       (2) INFORMATION FOR SEQ ID NO: 440:
 45
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 54 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 440:
 50
       Met Tyr Leu Cys Lys Thr Val Lys Val Leu Ile Cys Tyr Asp Trp Ile
        Leu Gly Leu Val Ser Ser Gly Gln His Trp Val Val Ser Leu Ser Tyr
  55
                     20
        Ser Ile Arg Val Tyr Pro Ala Met His Phe Thr Leu Cys Val His Ile
                                     40
  60
        Tyr Ser Lys Glu Pro Cys
```

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5	(2) INFORMATION FOR SEQ ID NO: 441:
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 42 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 441:
15	Met Thr Ala Leu Val Trp Arg Lys Gly Pro Asp Gly Gly Ser Arg Lys 1 5 10 15 Pro Ile Leu Leu Leu Phe Phe Phe Leu Pro Leu Ile Leu Cys Phe His
20	20 25 30 Ser Phe Ile His Ser Ser Asn Ile Cys Xaa 35 40
25	(2) INFORMATION FOR SEQ ID NO: 442: (i) SEQUENCE CHARACTERISTICS:
30	(A) LENGTH: 66 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 442:
	Met Phe Leu Thr Thr Trp Phe Leu Leu Ser Val Ala Trp Xaa Ala 1 5 10 15
35	Leu Thr Arg Ser Gly Arg Ser Cys Leu Pro Leu Val Gly Arg Pro Arg 20 25 30
40	Glu Gln Ser Pro Arg Thr His Cys Ala Ala Ser Ser Thr Lys Glu Arg 35 40 45 Asn Ser Asp Pro Gln Pro Ser Pro Pro Glu Val Val Gly Pro Leu Trp
	50 55 60 Ser Xaa
45	65
50	(2) INFORMATION FOR SEQ ID NO: 443: (i) SEQUENCE CHARACTERISTICS:
. 55	(A) LENGTH: 156 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 443:
	Met Lys Ala Ile Gly Ile Glu Pro Ser Leu Ala Thr Tyr His His Ile 1 5 10 15
60	Ile Arg Leu Phe Asp Gln Pro Gly Asp Pro Leu Lys Arg Ser Ser Phe

		20	25	30
	Ile Ile Tyr 35	Asp Ile Met	: Asn Glu Leu Met G 40	ly Lys Arg Phe Ser Pro 45
5	Lys Asp Pro 50	Asp Asp Asp	Lys Phe Phe Gln S 55	Ser Ala Met Ser Ile Cys 60
10	Ser Ser Leu 65	Arg Asp Leu 70		Sln Val His Gly Leu Leu 75 80
	Lys Thr Gly	Asp Asn Trp 85	Lys Phe Ile Gly 1	Pro Asp Gln His Arg Asn 95
15	Phe Tyr Tyr	Ser Lys Phe	e Phe Asp Leu Ile (105	Cys Leu Met Glu Gln Ile 110
20	Asp Val Thr		p Tyr Glu Asp Leu 120	Ile Pro Ser Ala Tyr Phe 125
20	Pro His Ser 130	Gln Thr Me	t Ile His Leu Leu 135	Gln Ala Leu Asp Val Ala 140
25	Asn Arg Leu 145	ı Glu Val Il 15	e Pro Lys Ile Trp 0	Glu Arg 155
30		SEQUENCE CH	Q ID NO: 444: HARACTERISTICS: STH: 57 amino acide E: amino acid	5
35	(xi	(D) TOP	OLOGY: linear DESCRIPTION: SEQ I	O NO: 444:
	Met His Ph 1	e Leu Phe Ar 5	rg Phe Ile Val Phe 10	Phe Tyr Leu Trp Gly Leu 15
40	Phe Thr Al	a Gln Arg Gl 20	ln Lys Lys Glu Glu 25	Ser Thr Glu Glu Val Lys 30
45	3	15	rg Pro Glu Asn Cys 40 ys Pro Leu Xaa	Ser Lys Thr Ser Lys Lys 45
	50		55	
50			EQ ID NO: 445:	
5 5		(A) LEN (B) TYI (D) TOI	HARACTERISTICS: WOTH: 416 amino acid PC: amino acid POLOGY: linear DESCRIPTION: SEQ 1	
60	Met Arg T	hr Leu Phe A 5	sn Leu Leu Trp Leu 10	Ala Leu Ala Cys Ser Pro 15

	Val I	His	Thr	Thr 20	Leu	Ser	Lys	Ser	Asp 25	Ala	Lys	Lys	Ala	Ala 30	Ser	Lys
5	Thr !	Leu	Leu 35	Glu	Lys	Ser	Gln	Phe 40	Ser	Asp	Lys	Pro	Val 45	Gln	Asp	Arg
10	Gly	Leu 50		Val	Thr	Asp	Leu 55	Lys	Ala	Glu	Ser	Val 60	Val	Leu	Glu	His
10	Arg 65	Ser	Tyr	Cys	Ser	Ala 70		Ala	Arg	Asp	Arg 75	His	Phe	Ala	Gly	Asp 80
15	Val	Leu	Gly	Tyr	Val 85		Pro	Trp	Asn	Ser 90		Gly	Tyr	Asp	Val 95	Thr
	Lys	Val	Phe	Gly 100		Lys	Phe	Thr	Gln 105		e Ser	Pro	Val	Trp 110	Leu	Gln
20	Leu	Lys	Arg		Gly	Arg	Glu	120		Glu	ı Val	Thr	Gly 125	Leu	His	Asp
25	Val	Asp 130		ı Gly	Trp	Met	139		(Va)	Arç	j Lys	140	Ala	Lys	Gly	Leu
23.	His 145	Ile	e Val	l Pro	Arg	J Let 150		ı Phe	e Glu	ı Ası	155	Thi	туг	Asp) Asi	Phe 160
30					16	5				17	0				17:	
	Val	Va:	l Gl	n Va 18		a Ly:	s As	n Gli	n Hi 18		e As	p Gl	y Phe	2 Va:	L Va)	l Glu
35	Val	Tr	p As: 19		n Le	u Le	u Se	r Gli 20		s Ar	g Va	1 Gl	y Let 20	ı Ile S	e Hi	s Met
40	Leu	Th. 21		s Le	u Al	a Gl	u Al 21		u Hi	s Gl	n Al	a Ar 22	g Le O	u Le	u Al	a Leu
	Lev 225		1 11	e Pr	o Pr	o Al 23		e Th	r Pr	o G1	y Th 23	r As	p Gl	n Le	u Gl	y Met 240
45	Phe	e Th	ır Hi	s Ly	rs G1 24		ie Gl	u Gl	n Le	25 25		o Va	l Le	nu As	p G) 25	y Phe
	Sei	r Le	eu Me	et Th 26		n As	p T)	nc Se	r Tì 20		la Hi	is Gi	n Pr	o Gl 27	y Pi 0	o Asn
50	Ala	a Pı		eu Se 75	er T	rp Va	al A		la Co 30	ys Va	al G	ln Va	1 Le 28	eu As 35	p Pi	ro Lys
55	Se		ys T: 90	rp A	rg S	er Ly		le Le 95	eu L	eu G	ly L	eu As	sn Pl 00	ne Ty	r G	ly Met
J J	As 30		yr A	la T	hr S		ys A 10	sp A	la A	rg G	lu P	ro V	al Va	al G	ly A	la Arg 320
60	ту	r I	le G	ln T		eu L 25	ys A	sp H	is A		ro A 30	rg M	et V	al T	rp A 3	sp Ser 35

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	Gln	Xaa	Ser	Glu 340	His	Phe	Phe	Glu	Tyr 345	Lys	Lys	Ser	Arg	Ser 350	Gly	Arg	
5	His	Val	Val 355	Phe	Туг	Pro	Thr	Leu 360	Lys	Ser	Leu	Gln	Val 365	Arg	Leu	Glu	
10	Leu	Ala 370	Arg	Glu	Leu	Gly	Val 375	Gly	Val	Ser	Ile	Trp 380	Glu	Leu	Ala	Arg	í
i O	Ala 385	_	Thr	Thr	Ser	Thr 390	Thr	Cys	Ser	Arg	Trp 395	Ala	Leu	Arg	Pro	Pro 400	, }
15	Arg	Trp	Thr	Cys	Ser 405		Leu	Ser	His	Gly 410		Ser	Glu	Gln	Val 415	Xaa	ı
20																	
25	(2)	INF	(i)	SEQU	JENCE (A) I (B) ' (D) '	E CHI	RAC TH: : am LOGY	TERIS 64 au ino a	STICS mino acid near	S: acid		o: 4	1 6:				
30		t Ala	a Pro	Gly	/ Pro		ı Se	r Ala	a Thi	Glr 10		a Val	l Val	l Ile	His	Th	r
35				20	0				2	5				ı Val 30 u Arg)		
40			3 s Ly	5				. 40 u Xa	0				49 1 Pr				
45						- ~-			447								
	(2	() IN			UENC	TE CF	iarac	TERI	STIC	:S:							
50			(x:	i) SI	(B)	TYP	E: au	206 mino Y: 1: RIPT	acio inea	d r		10: 4	147:				
55	Me	et Le 1	eu Gl	ly Al	ia Ly	/s P1 5	o H	is Tı	np Le		ro G1 L0	y Pı	co Le	eu Hi		er P 15	ro
60	G:	ly Le	eu P		eu Va 20	al Le	eu Va	al Le		eu A] 25	la Le	eu G	ly Al	la Gl	y T: 30	np A	la

•	Gln Glu Gly Ser Glu Pro Val Leu Leu Glu Gly Glu Cys Leu Val Val 35 40 45
5	Cys Glu Pro Gly Arg Ala Ala Ala Gly Gly Pro Gly Gly Ala Ala Leu 50 55 60
	Gly Glu Ala Pro Pro Gly Arg Val Ala Phe Ala Ala Val Arg Ser Xaa 65 70 75 80
10	His His Glu Pro Ala Gly Glu Thr Gly Asn Gly Thr Xaa Gly Ala Ile 85 90 95
15	Tyr Phe Asp Gln Val Leu Val Asn Glu Gly Gly Gly Phe Asp Arg Ala 100 105 110
15	Ser Gly Ser Phe Val Ala Pro Val Arg Gly Val Tyr Ser Phe Arg Phe 115 120 125
20	His Val Val Lys Val Tyr Asn Arg Gln Thr Val Gln Val Ser Leu Met 130 135 140
	Leu Asn Thr Trp Pro Val Ile Ser Ala Phe Ala Asn Asp Pro Asp Val 145 150 155 160
25	Thr Arg Glu Ala Ala Thr Ser Ser Val Leu Leu Pro Leu Asp Pro Gly 165 170 175
30	Asp Arg Val Ser Leu Arg Leu Arg Gly Asn Leu Leu Gly Gly Trp 180 185 190
30	Lys Tyr Ser Ser Phe Ser Gly Phe Leu Ile Phe Pro Leu Xaa 195 200 205
35 -	(2) INFORMATION FOR SEQ ID NO: 448:
	(i) SEQUENCE CHARACTERISTICS:
40	(A) LENGTH: 62 amino acids (B) TYPE: amino acid
40	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 448:
• •	
45	Met Ser Ser Leu Leu Ser Ala Gly Leu Gln Ala Ser Leu Cys Gly Lys 1 5 10 15
	Xaa Leu Trp Ala Ser Thr Trp Tyr Leu Val Cys Cys Leu Leu Pro Phe 20 25 30
50	35 40 45
	Asn Leu Lys Ser Tyr Cys Gly Leu Ser Thr Ile Glu Ile Xaa 50 55 60
55	
	(2) INFORMATION FOR SEQ ID NO: 449:
60	(i) SEQUENCE CHARACTERISTICS:

•	(A) LENGTH: 316 amino acids (B) TYPE: amino acid
	(D) TOPOLOGY: linear
_	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 449:
5	Met Ser Thr Lys Lys Leu Cys Ile Val Gly Gly Ile Leu Leu Val Phe 1 5 10 15
10	Gln Ile Ile Ala Phe Leu Val Gly Gly Leu Ile Ala Pro Gly Pro Thr 20 25 30
	Thr Ala Val Ser Tyr Met Ser Val Lys Cys Val Asp Ala Arg Lys Asn 35 40 45
15	His His Lys Thr Lys Trp Phe Val Pro Trp Gly Pro Asn His Cys Asp 50 55 60
20	Lys Ile Arg Asp Ile Glu Glu Ala Ile Pro Arg Glu Ile Glu Ala Asn 65 70 75 80
20	Asp Ile Val Phe Ser Val His Ile Pro Leu Pro His Met Glu Met Ser 85 90 95
25	Pro Trp Phe Gln Phe Met Xaa Phe Ile Leu Gln Leu Asp Ile Ala Phe 100 105 110
	Lys Leu Asn Asn Gln Ile Arg Glu Asn Ala Glu Val Ser Met Asp Val 115 120 125
30	Ser Leu Ala Tyr Arg Asp Asp Ala Phe Ala Glu Trp Thr Glu Met Ala 130 135 140
35	His Glu Arg Val Pro Arg Lys Leu Lys Cys Thr Phe Thr Ser Pro Lys 145 150 155 160
33	Thr Pro Glu His Gly Gly Pro Val Thr Met Asn Val Met Ser Phe Leu 165 170 175
40	Ser Trp Lys Leu Gly Leu Trp Pro Met Lys Phe Tyr Leu Leu Asn Ile 180 185 190
٠.	Arg Leu Pro Val Asn Glu Lys Lys Lys Ile Asn Val Gly Ile Gly Glu 195 200 205
45	Ile Lys Asp Ile Arg Leu Val Gly Ile His Gln Asn Gly Gly Phe Thr 210 215 220
50	Lys Val Trp Phe Ala Met Lys Thr Phe Leu Thr Pro Ser Ile Phe Ile 225 230 235 240
	Ile Met Val Trp Tyr Trp Arg Arg Ile Thr Met Met Ser Arg Pro Pro 245 250 255
55	·
	Ile Asn Ile Pro Val Glu Trp Phe Ser Ile Gly Phe Asp Trp Thr Trp 275 280 285
60) Met Leu Leu Phe Gly Asp Ile Arg Gln Ala Ser Ser Met Xaa Cys Ph

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300
                             295
         290
     Xaa Pro Ser Gly Ser Ser Ser Val Ala Ser Thr Xaa
                         310
5
      (2) INFORMATION FOR SEQ ID NO: 450:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 24 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 450:
15
      Met Leu Ala Leu Leu Gly Leu Leu Ala Gly Thr Glu His Pro Pro Gly
                        5
      Pro Gln Gly Pro Gly Pro Ser Xaa
20
                   20
      (2) INFORMATION FOR SEQ ID NO: 451:
25
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 10 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 451:
30
      Met Pro Ser Gly Ala Cys Cys Ser Pro Xaa
                        5
        1
 35
       (2) INFORMATION FOR SEQ ID NO: 452:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 26 amino acids
 40
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 452:
       Met Leu Pro Ala Leu Ser Thr Val Leu Leu Pro Thr Pro Ser Leu Cys
 45
                        5
       Ser Gly Asn Pro Arg Glu Gly Trp Ala Xaa
                    20
 50
        (2) INFORMATION FOR SEQ ID NO: 453:
               (i) SEQUENCE CHARACTERISTICS:
  55
                      (A) LENGTH: 172 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 453:
  60
```

	Met 1	Tyr	Ser	Leu	His 5	Ser	Trp	Val	Gly	Leu 10	Ile	Ala	Val	Ile	Cys 15	Tyr
5	Leu	Leu	Gln	Leu 20	Leu	Ser	Gly	Phe	Ser 25	Val	Phe	Leu	Leu	Pro 30	Trp	Ala
	Pro	Leu	Ser 35	Leu	Arg	Ala	Phe	Leu 40	Met	Pro	Ile	His	Val 45	Tyr	Ser	Gly
10	Ile	Val 50	Ile	Phe	Gly	Thr	Val 55	Ile	Ala	Thr	Ala	Leu 60	Met	Gly	Leu	Thr
15	Glu 65	Lys	Leu	Ile	Phe	Ser 70	Leu	Arg	Asp	Pro	Ala 75	Tyr	Ser	Thr	Phe	Pro 80
	Pro	Glu	Gly	Val	Phe 85	Val	Asn	Thr	Leu	90	Leu	Leu	Ile	Leu	Val 95	Phe
20				100					105					110		
	Lys	Glu	Pro 115		Ser	Thr	Ile	Leu 120	His	Pro	Asn	Gly	Gly 125	Thr	Glu	Gln
25	Gly	Ala 130		Gly	Ser	Met	Pro 135		Tyr	Ser	Gly	Asn 140	Asn	Met	Asp	Lys
30	Ser 145		Ser	Glu	. Leu	Asn 150		Glu	Val	Ala	Ala 155		Lys	Arg	Asn	Leu 160
	Ala	Leu	Asp	Glu	Ala 165	-	Gln	Arg	Ser	Thr 170		Xaa				
35	(2)	INF	ORMA	TION	FOR	SEÇ) ID	NO:	454:							
40					JENCE (A) I (B) ' (D) '	ENG TYPE TOPO	TH: : : am LOGY	96 ar ino a : li	mino acid near	acio		o: 4 5	i4:			
45		: Phe	e His	s Val	l Leu		: Ala	Glr	val	Thr 10		Val	. Ile	Ile	Thr 15	Thr
	Va:	l Sei	r Va	l Let 20		Phe	e Ası	Phe	Arg 25	_	Ser	Leu	Glu	Phe 30		. Leu
50	Gli	u Ala	а Ха 3	_	r Val	l Xaa	a Let	ı Sei 40		Phe	e Ile	е Туг	Asr 49		. Ser	Lys
55	Pr	6 Gl		l Pr	o Glu	1 Ту	r Ala) Arq	g Gli	n Glu	Arg 60		e Arg	j Asi	Leu
<i></i>	Se 6		y As	n Le	u Trj	9 Gl		g Se:	r Se	r Gly	y As _] 7:		y Glu	ı Glu	ı Let	Glu 80
60	Ar	g Le	u Th	r Ly	s Pr		s Se	r As	p Gl	u Se:		o Gl	ı Ası	Thu	Phe 99	Xaa S

5	
	(2) INFORMATION FOR SEQ ID NO: 455:
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 171 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 455:
15	Met Arg Gly Pro Ala Gln Ala Lys Leu Leu Pro Gly Ser Ala Ile Gln 1 5 10 15
	Ala Leu Val Gly Leu Ala Arg Pro Leu Val Leu Ala Leu Leu Val 20 25 30
20	Ser Ala Ala Leu Ser Ser Val Val Ser Arg Thr Asp Ser Pro Ser Pro 35 40 45
25	Thr Val Leu Asn Ser His Ile Ser Thr Pro Asn Val Asn Ala Leu Thr 50 55 60
	His Glu Asn Gln Thr Lys Pro Ser Ile Ser Gln Ile Ser Thr Thr Leu 65 70 75 80
30	Pro Pro Thr Thr Ser Thr Lys Lys Ser Gly Gly Ala Ser Val Val Pro 85 90 95
35	His Pro Ser Pro Thr Pro Leu Ser Gln Glu Glu Ala Asp Asn Asn Glu 100 105 110
33	Asp Pro Ser Ile Glu Glu Glu Asp Leu Leu Met Leu Asn Ser Ser Pro 115 120 125
40	Ser Thr Ala Lys Asp Thr Leu Asp Asn Gly Asp Tyr Gly Glu Pro Asp 130 135 140
	Tyr Asp Trp Thr Thr Gly Pro Arg Asp Asp Asp Glu Ser Asp Xaa His 145 150 155 160
45	Leu Gly Arg Lys Gln Gly Leu His Gly Asn Xaa 165 170
50	(2) INFORMATION FOR SEQ ID NO: 456:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids
55	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 456:
	Met Lys Ala Ser Gln Cys Cys Cys Cys Leu Ser His Leu Leu Ala Ser
	1 5 10 15

	Val Leu Leu Leu Leu Leu Pro Glu Leu Ser Gly Xaa Leu Xaa Val 20 25 30
5	Leu Leu Gln Ala Ala Glu Ala Ala Pro Gly Xaa Gly Pro Pro Asp Pro 35 40 45
	Arg Pro Gly His Tyr Arg Arg Cys His Arg Ala Leu Thr Pro Ala Gln 50 55 60
10	Gln Pro Gly Arg Gly Leu Ala Glu Ala Ala Gly Ala Ala Gly Leu Arg 65 70 75 80
15	Gly Arg Gln Trp Gln Gln Pro Cys Gly Arg Ala Xaa 85 90
	(2) INFORMATION FOR SEQ ID NO: 457:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 206 amino acids (B) TYPE: amino acid
25	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 457: Ile Ser Val Leu Xaa Tyr Pro His Cys Val Val His Glu Leu Pro Glu
	1 5 10 15 Leu Thr Ala Glu Ser Leu Glu Ala Gly Asp Ser Asn Gln Phe Cys Trp
30	20 25 30
	Arg Asn Leu Phe Ser Cys Ile Asn Leu Leu Arg Ile Leu Asn Lys Leu 35 40 45
35	Thr Lys Trp Lys His Ser Arg Thr Met Met Leu Val Val Phe Lys Ser 50 55 60
40	Ala Pro Ile Leu Lys Arg Ala Leu Lys Val Lys Gln Ala Met Met Gln 65 70 75 80
٠.	Leu Tyr Val Leu Lys Leu Leu Lys Val Gln Thr Lys Tyr Leu Gly Arg 85 90 95
45	Gln Trp Arg Lys Ser Asn Met Lys Thr Met Ser Ala Ile Tyr Gln Lys 100 105 110
	Val Arg His Arg Leu Asn Asp Asp Trp Ala Tyr Gly Asn Asp Leu Asp 115 120 125
50	Ala Arg Pro Trp Asp Phe Gln Ala Glu Glu Cys Ala Leu Arg Ala Asn 130 135 140
55	Ile Glu Arg Phe Asn Ala Arg Arg Tyr Asp Arg Ala His Ser Asn Pro 145 150 150 160
	Asp Phe Leu Pro Val Asp Asn Cys Leu Gln Ser Val Leu Gly Gln Arg 165 170 175
60	Val Asp Leu Pro Glu Asp Phe Gln Met Asn Tyr Asp Leu Trp Leu Glu

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Arg Glu Val Phe Ser Lys Pro Ile Ser Trp Glu Glu Leu Leu
195 200 205

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5		
	(2) INFORMATION FOR SEQ ID NO: 458:	
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 317 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 458:	
15	Met Ala Pro Pro Ala Pro Gly Pro Ala Ser Gly Gly Ser Gly Glu Val	L
20	Asp Glu Leu Phe Asp Val Lys Asn Ala Phe Tyr Ile Gly Ser Tyr Gli 20 25 30	a
20	Gln Cys Ile Asn Glu Ala Xaa Xaa Val Lys Leu Ser Ser Pro Glu Ar 35 40 45	g
25	Asp Val Glu Arg Asp Val Phe Leu Tyr Arg Ala Tyr Leu Ala Gln Ar 50 55 60	g
	Lys Phe Gly Val Val Leu Asp Glu Ile Lys Pro Ser Ser Ala Pro Gl 65 70 75 8	u 10
30	Leu Gln Ala Val Arg Met Phe Ala Asp Tyr Leu Ala His Glu Ser Ar 85 90 95	g
35	Arg Asp Ser Ile Val Ala Glu Leu Asp Arg Glu Met Ser Arg Ser Xa 100 105 110	ıa
33	Asp Val Thr Asn Thr Thr Phe Leu Leu Met Ala Ala Ser Ile Tyr Le 115 120 125	eu
40	His Asp Gln Asn Pro Asp Ala Ala Leu Arg Ala Leu His Gln Gly A 130 135 140	gz
,	Ser Leu Glu Cys Thr Ala Met Thr Val Gln Ile Leu Leu Lys Leu A 145 150 155 1	sp 60
45	Arg Leu Asp Leu Ala Arg Lys Glu Leu Lys Arg Met Gln Asp Leu A 165 170 175	sp
50	Glu Asp Ala Thr Leu Thr Gln Leu Ala Thr Ala Trp Val Ser Leu A 180 185 190	la
30	Thr Gly Gly Glu Lys Leu Gln Asp Ala Tyr Tyr Ile Phe Gln Glu E 195 200 205	let
55	Ala Asp Lys Cys Ser Pro Thr Leu Leu Leu Leu Asn Gly Gln Ala 2 210 215 220	l la
	Cys His Met Ala Gln Gly Arg Trp Glu Ala Ala Glu Gly Leu Leu C 225 230 235	31r 240

60 Glu Ala Leu Asp Lys Asp Ser Gly Tyr Pro Glu Thr Leu Val Asn Leu

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-	245 250 255	
_	Ile Val Leu Ser Gln His Leu Gly Lys Pro Pro Glu Val Thr Asn Arg 260 265 270	
5	Tyr Leu Ser Gln Leu Lys Asp Ala His Arg Ser His Pro Phe Ile Lys 275 280 285	
10	Glu Tyr Gln Ala Lys Glu Asn Asp Phe Asp Arg Leu Val Leu Gln Tyr 290 295 300	
	Ala Pro Ser Ala Glu Ala Gly Pro Glu Leu Ser Gly Pro 305 310 315	
15	AS9	
20	(2) INFORMATION FOR SEQ ID NO: 459: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 261 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 459:	
25	Arg Asp Val Glu Arg Asp Val Phe Leu Tyr Arg Ala Tyr Leu Ala Gln 1 5 10 15	
30	Arg Lys Phe Gly Val Val Leu Asp Glu Ile Lys Pro Ser Ser Ala Pro 20 25 30 Glu Leu Gln Ala Val Arg Met Phe Ala Asp Tyr Leu Ala His Glu Ser	
35	35 40 45 - Arg Arg Asp Ser Ile Val Ala Glu Leu Asp Arg Glu Met Ser Arg Ser 50 55 60	
	Xaa Asp Val Thr Asn Thr Thr Phe Leu Leu Met Ala Ala Ser Ile Tyr 65 70 75 86	•
40	Leu His Asp Gln Asn Pro Asp Ala Ala Leu Arg Ala Leu His Gln Gl 85 90 95	
45	Asp Ser Leu Glu Cys Thr Ala Met Thr Val Gln Ile Leu Leu Lys Leu 100 105 110 Asp Arg Leu Asp Leu Ala Arg Lys Glu Leu Lys Arg Met Gln Asp Leu 115 120 125	
50	Asp Glu Asp Ala Thr Leu Thr Gln Leu Ala Thr Ala Trp Val Ser Le 130 135 140	
	Ala Thr Gly Gly Glu Lys Leu Gln Asp Ala Tyr Tyr Ile Phe Gln Gl 145 150 155 16	1u 50
55	Met Ala Asp Lys Cys Ser Pro Thr Leu Leu Leu Leu Asn Gly Gln A 165 170 175	la
60	Ala Cys His Met Ala Gln Gly Arg Trp Glu Ala Ala Glu Gly Leu L 180 185 190	eu

•	Gln Glu Ala Leu Asp Lys Asp Ser Gly Tyr Pro Glu Thr Leu Val Asn 195 200 205
5	Leu Ile Val Leu Ser Gln His Leu Gly Lys Pro Pro Glu Val Thr Asn 210 215 220
	Arg Tyr Leu Ser Gln Leu Lys Asp Ala His Arg Ser His Pro Phe Ile 225 230 235 240
10	Lys Glu Tyr Gln Ala Lys Glu Asn Asp Phe Asp Arg Leu Val Leu Gln 245 250 255
15	Tyr Ala Pro Ser Ala 260
	(2) INFORMATION FOR SEQ ID NO: 460:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 156 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 460:
	Met Lys Ala Ile Gly Ile Glu Pro Ser Leu Ala Thr Tyr His His Ile 1 5 10 15
30	Ile Arg Leu Phe Asp Gln Pro Gly Asp Pro Leu Lys Arg Ser Ser Phe 20 25 30
	Ile Ile Tyr Asp Ile Met Asn Glu Leu Met Gly Lys Arg Phe Ser Pro 35 40 45
35	Lys Asp Pro Asp Asp Asp Lys Phe Phe Gln Ser Ala Met Ser Ile Cys 50 55 60
40	Ser Ser Leu Arg Asp Leu Glu Leu Ala Tyr Gln Val His Gly Leu Leu 65 70 75 80
40	Lys Thr Gly Asp Asn Trp Lys Phe Ile Gly Pro Asp Gln His Arg Asn 85 90 95
45	Phe Tyr Tyr Ser Lys Phe Phe Asp Leu Ile Cys Leu Met Glu Gln Ile 100 105 110
	Asp Val Thr Leu Lys Trp Tyr Glu Asp Leu Ile Pro Ser Ala Tyr Phe 115 120 125
50	Pro His Ser Gln Thr Met Ile His Leu Leu Gln Ala Leu Asp Val Ala 130 135 140
55	Asn Arg Leu Glu Val Ile Pro Lys Ile Trp Glu Arg 145 150 155
	(2) INFORMATION FOR SEQ ID NO: 461:

(i) SEQUENCE CHARACTERISTICS:

	(A) LENGTH: 176 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 461:
5	Lys Asp Ser Lys Glu Tyr Gly His Thr Phe Arg Ser Asp Leu Arg Glu 1 5 10 15
10	Glu Ile Leu Met Leu Met Ala Arg Asp Lys His Pro Pro Glu Leu Gln 20 25 30
	Val Ala Phe Ala Asp Cys Ala Ala Asp Ile Lys Ser Ala Tyr Glu Ser 35 40 45
15	Gln Pro Ile Arg Gln Thr Ala Gln Asp Trp Pro Ala Thr Ser Leu Asn 50 55 60
20	Cys Ile Ala Ile Leu Phe Leu Arg Ala Gly Arg Thr Gln Glu Ala Trp 65 70 75 80
20	Lys Met Leu Gly Leu Phe Arg Lys His Asn Lys Ile Pro Arg Ser Glu 85 90 95
25	Leu Leu Asn Glu Leu Met Asp Ser Ala Lys Val Ser Asn Ser Pro Ser 100 105 110
	Gln Ala Ile Glu Val Val Glu Leu Ala Ser Ala Phe Ser Leu Pro Ile 115 120 125
30	Cys Glu Gly Leu Thr Gln Arg Val Met Ser Asp Phe Ala Ile Asn Gln 130 135 140
35	Glu Gln Lys Glu Ala Leu Ser Asn Leu Thr Ala Leu Thr Ser Asp Ser 145 150 155 160
	Asp Thr Asp Ser Ser Ser Asp Ser Asp Ser Asp Thr Ser Glu Gly Lys 165 170 175
40	
45	(2) INFORMATION FOR SEQ ID NO: 462:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 324 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 462:
	Met Ser Ser Asp Asn Glu Ser Asp Ile Glu Asp Glu Asp Leu Lys Leu 1 5 10 15
55	Glu Leu Arg Arg Leu Arg Asp Lys His Leu Lys Glu Ile Gln Asp Leu 20 25 30
60	Gln Ser Arg Gln Lys His Glu Ile Glu Ser Leu Tyr Thr Lys Leu Gly 35 40 45

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	Lys Val Pro Pro Ala Val Ile Ile Pro Pro Ala Ala Pro Leu Ser Gly 50 55 60
5	Arg Arg Arg Pro Thr Lys Ser Lys Gly Ser Lys Ser Ser Arg Ser 65 70 75 80
	Ser Ser Leu Gly Asn Lys Ser Pro Gln Leu Ser Gly Asn Leu Ser Gly 85 90 95
10	Gln Ser Ala Ala Ser Val Leu His Pro Gln Gln Thr Leu His Pro Pro 100 105 110
15	Gly Asn Ile Pro Glu Ser Gly Gln Asn Gln Leu Leu Gln Pro Leu Lys 115 120 125
13	Pro Ser Pro Ser Ser Asp Asn Leu Tyr Ser Ala Phe Thr Ser Asp Gly 130 135 140
20	Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly Gln Gly Thr Ser Ser 145 150 155 160
	Thr Asn Thr Val Gly Ala Thr Val Asn Ser Gln Ala Ala Gln Ala Gln 165 170 175
25	Pro Pro Ala Met Thr Ser Ser Arg Lys Gly Thr Phe Thr Asp Asp Leu 180 185 190
30	His Lys Leu Val Asp Asn Trp Ala Arg Asp Ala Met Asn Leu Ser Gly 195 200 205
	Arg Arg Gly Ser Lys Gly His Met Asn Tyr Glu Gly Pro Gly Met Ala 210 215 220
35	Arg Lys Phe Ser Ala Pro Gly Gln Leu Cys Ile Ser Met Thr Ser Asn 225 230 235 240
	Leu Gly Gly Ser Ala Pro Ile Ser Ala Ala Ser Ala Thr Ser Leu Gly 245 250 255
40	His Phe Thr Lys Ser Met Cys Pro Pro Gln Gln Tyr Gly Phe Pro Ala 260 265 270
45	Thr Pro Phe Gly Ala Gln Trp Ser Gly Thr Gly Gly Pro Ala Pro Gli 275 280 285
	Pro Leu Gly Gln Phe Gln Pro Val Gly Thr Ala Ser Leu Gln Asn Ph 290 295 300
50	Asn Ile Ser Asn Leu Gln Lys Ser Ile Ser Asn Pro Pro Gly Ser As 305 310 315 32
	Leu Arg Thr Thr
55	

(2) INFORMATION FOR SEQ ID NO: 463:

60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 133 amino acids

•	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 463:													
5	Ile Gln Asp Leu Gln Ser Arg Gln Lys His Glu Ile Glu Ser Leu Tyr 1 5 10 15													
10	Thr Lys Leu Gly Lys Val Pro Pro Ala Val Ile Ile Pro Pro Ala Ala 20 25 30													
10	Pro Leu Ser Gly Arg Arg Arg Pro Thr Lys Ser Lys Gly Ser Lys 35 40 45													
15	Ser Ser Arg Ser Ser Ser Leu Gly Asn Lys Ser Pro Gln Leu Ser Gly 50 55 60													
	Asn Leu Ser Gly Gln Ser Ala Ala Ser Val Leu His Pro Gln Gln Thr 65 .70 .75 .80													
20	Leu His Pro Pro Gly Asn Ile Pro Glu Ser Gly Gln Asn Gln Leu Leu 85 90 95													
25	Gln Pro Leu Lys Pro Ser Pro Ser Ser Asp Asn Leu Tyr Ser Ala Phe 100 105 110													
23	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly Gln 115 120 125													
30	Gly Thr Ser Ser Thr 130													
35	(2) INFORMATION FOR SEQ ID NO: 464: (i) SEQUENCE CHARACTERISTICS:													
	(A) LENGTH: 53 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear													
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 464:													
٠.	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly Glr 1 5 10 15													
45	Gly Thr Ser Ser Thr Asn Thr Val Gly Ala Thr Val Asn Ser Gln Ala 20 25 30													
50	Ala Gln Ala Gln Pro Pro Ala Met Thr Ser Ser Arg Lys Gly Thr Ph 35 40 45													
50	Thr Asp Asp Leu His													
55	(2) INFORMATION FOR SEQ ID NO: 465:													
	(i) SEQUENCE CHARACTERISTICS:													
60	(A) LENGTH: 48 amino acids (B) TYPE: amino acid													

•	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 465:
5	Lys Gly His Met Asn Tyr Glu Gly Pro Gly Met Ala Arg Lys Phe Ser 1 5 10 15
	Ala Pro Gly Gln Leu Cys Ile Ser Met Thr Ser Asn Leu Gly Gly Ser 20 25 30
10	Ala Pro Ile Ser Ala Ala Ser Ala Thr Ser Leu Gly His Phe Thr Lys 35 40 45
15	
	(2) INFORMATION FOR SEQ ID NO: 466:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 466:
23	Gln Pro Leu Lys Pro Ser Pro Ser Ser Asp Asn Leu Tyr Ser Ala Phe 1 5 10 15
30	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly 20 25 30 .
35	(2) INFORMATION FOR SEQ ID NO: 467:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 57 amino acids
	(B) TYPE: amino acid (D) TOPOLOGY: linear
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 467:
• •	Val Arg Val Ala Ala Ala Glu Ser Met Xaa Leu Leu Leu Glu Cys Ala 1 5 10 15
45	Xaa Val Arg Gly Pro Glu Tyr Leu Thr Gln Met Trp His Phe Met Cys 20 25 30
	Asp Ala Leu Ile Lys Ala Ile Gly Thr Glu Pro Asp Ser Asp Val Let 35 40 45
50	Ser Glu Ile Met His Ser Phe Ala Lys 50 55
55	(2) INFORMATION FOR SEQ ID NO: 468:
-	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 85 amino acids (B) TYPE: amino acid
60	J (U)

	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 468:
5	Met Glu Ile Asn Asn Gln Asn Cys Phe Ile Val Ile Asp Leu Val Arg 1 5 10 15
	Thr Val Met Glu Asn Gly Val Glu Gly Leu Leu Ile Phe Gly Ala Phe 20 25 30
10	Leu Pro Glu Ser Trp Leu Ile Gly Val Arg Cys Ser Ser Glu Pro Pro 35 40 45
	Lys Ala Leu Leu Leu Ile Leu Ala His Ser Gln Lys Arg Arg Leu Asp 50 55 60
15	Gly Trp Ser Phe Ile Arg His Leu Arg Val His Tyr Cys Val Ser Leu 65 70 75 80
20	Thr Ile His Phe Ser 85
25	(2) INFORMATION FOR SEQ ID NO: 469:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids (B) TYPE: amino acid
30	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 469:
	Gln Asp Lys His Ala Glu Glu Val Arg Lys Asn Lys Glu Leu Lys Glu 1 5 10 15
35	Glu Ala Ser Arg 20
40	(2) INFORMATION FOR SEQ ID NO: 470:
٠.	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids
45	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 470:
	Gln Gln Asp Leu Ser Pro Trp Ala Ala Pro Val Gly Cys Pro Leu Xaa 1 5 10 15
50	Xaa Ala Ser Xaa Thr Cys His Xaa Leu Pro Leu Ser Gly Cys Leu Arg 20 25 30
55	Arg Gln Ser Xaa Ser Leu Pro Val Val Ala Xaa Leu Cys Phe Trp Phe 35 40 45
	Ser Cys Pro Leu Ala Ser Leu Phe Val Pro Gly Gln Pro Cys Val Thr 50 55 60
60	Cys Pro Phe Pro Ser Leu Pro Phe Gln Asp Lys His Ala Glu Glu Val

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80
                          70
     55
     Arg Lys Asn Lys Glu Leu Lys Glu Glu Ala Ser Arg
                      35
5
     (2) DESCRIPTION FOR SEQ ID NO: 471:
             ::) SEQUENCE CHARACTERISTICS:
10
                   (A) LENGTH: 37 amino acids
                    (2) TYPE: amino acid
                    (D) TOPOLOGY: linear
              MI; SEQUENCE DESCRIPTION: SEQ ID NO: 471:
15
      Pro The Arg Cys Cys Thr Thr Gln Pro Cys Arg Ser Ser Ala Arg Arg
      Pro Cys Trp Val Pro Met Val Pro Ser Pro Glu Gly Arg Glu Kaa Gln
20
      Pro Thr Cys Pro Ser
25
       (2) DECEMBERCH FOR SEQ ID NO: 472:
              (1) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 363 amino acids
 30
                     (3) TYPE: amino acid
                     (D) TOPOLOGY: linear
               'xi) SEQUENCE DESCRIPTION: SEQ ID NO: 472:
       Met Lys Arg Ser Leu Asn Glu Asn Ser Ala Arg Ser Thr Ala Gly Cys
 35
                                          10
       Leu Pro Val Pro Leu Phe Asn Gln Lys Lys Arg Asn Arg Gln Pro Leu
                                   25
 40
       Thr Ser Asm Pro Leu Lys Asp Asp Ser Gly Ile Ser Thr Pro Ser Asp
                                    40
       Asn Tyr Asp Phe Pro Pro Leu Pro Thr Asp Trp Ala Trp Glu Ala Val
  45
        Asn Pro Glu Zaa Ala Pro Val Met Lys Thr Val Asp Thr Gly Gln Ile
        Pro His Ser Val Ser Arg Pro Leu Arg Ser Gln Asp Ser Val Phe Asn
  50
        Ser Ile Glm Ser Asn Thr Gly Arg Ser Gln Gly Gly Trp Ser Tyr Arg
                                        105
  55
        Asp Sly Asm Lys Asm Thr Ser Leu Lys Thr Trp Xaa Lys Asm Asp Phe
                                    120
         Lys Pro Gln Cys Lys Arg Thr Asn Leu Val Ala Asn Asp Gly Lys Asn
                                135
   60
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•	Ser (Cys	Pro	Met	Ser	Ser 150	Gly	Ala	Gln	Gln	Gln 155	Lys	Gln	Leu	Arg	Thr 160
5	Pro (Glu	Pro	Pro	Asn 165	Leu	Ser	Arg	Asn	Lys 170	Glu	Thr	Glu	Leu	Leu 175	Arg
	Gln '	Thr	His	Ser 180		Lys	Ile	Ser	Gly 185	Cys	Thr	Met	Arg	Gly 190	Leu	Asp
10	Lys	Asn	Ser 195	Ala	Leu	Gln	Thr	Leu 200	Lys	Pro	Asn	Phe	Gln 205	Gln	Asn	Gln
15	Tyr	Lys 210	Xaa	Gln	Met	Leu	Asp 215		Ile	Pro	Glu	Asp 220	Asn	Thr	Leu	Lys
	Glu 225	Thr	Ser	Leu	Tyr	Gln 230		Gln	Phẹ	Lys	Glu 235	Lys	Ala	Ser	Ser	Leu 240
20	Arg	Ile	Ile	Ser	Ala 245		Ile	e Glu	Ser	Met 250		Tyr	Trp	Arg	Glu 255	His
25	Ala	Gln	Lys	Thr 260		. Leu	. Lev	. Phe	265		Leu	Ala	. Val	. Leu 270	Asp	Ser
23	Ala	Val	Thr 275		Gly	, Pro	туг	280		Ly	s Thr	Phe	289	Met	: Arg	Asp
30		290)				29	5				300)			ı Leu
	Pro 305		j Lei	ı Il	e Ar	31		g Va	l Hi	s Ar	g Cy: 31	s Vai	l Gly	y Ası	а Туг	Asp 320
35	Gln	Ly	s Ly:	s As	n Il 32		e Gl	n Cy:	s Va	1 Se 33		l Ar	g Pr	o Ala	331	r Val 5
40	Ser	Gl	u Gl	n Ly 34		r Ph	e Gl	n Al	a Ph 34		1 Ly	s Il	e Al	a As 35	p Va O	l Glu
	Met	: Gl	n Ty 35		r Il	e As	n Va	1 Me 36		n Gl	u Th	r				
45	(2)) IN	FORM	DITA	N FC	OR SE	EQ II	NO:	473	:						
50			(i)	SE((A)	LEN	GTH:	TERI 45 a mino	amin	o ac	ids					
20			(x	i) S	(D)	TOP	OLOG	Y: 1 RIPT	inea	r	ID 1	NO: 4	473:			
55		r G! 1	ln A	sp S	er Va	al Pi	he A	sn Se	er I		ln So 10	er A	sn T	hr G	ly A	rg Ser 15
	Gl	n G	ly G		rp S 20	er T	yr A	rg A		ly A 25	sn L	ys A	sn T	hr S	er L 30	eu Lys
60	Th	r T	rp X	aa L	ys A	sn A	sp P	he L	ys P	ro G	ln C	ys L	ys A	rg		

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40 45 35 (2) INFORMATION FOR SEQ ID NO: 474: 5 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 36 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 474: Asn Lys Glu Thr Glu Leu Leu Arg Gln Thr His Ser Ser Lys Ile Ser 15 Gly Cys Thr Met Arg Gly Leu Asp Lys Asn Ser Ala Leu Gln Thr Leu 25 Lys Pro Asn Phe 20 35 (2) INFORMATION FOR SEQ ID NO: 475: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 49 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 475: 30 Ser Ser Leu Arg Ile Ile Ser Ala Val Ile Glu Ser Met Lys Tyr Trp 1 Arg Glu His Ala Gln Lys Thr Val Leu Leu Phe Glu Val Leu Ala Val 35 25 20 Leu Asp Ser Ala Val Thr Pro Gly Pro Tyr Tyr Ser Lys Thr Phe Leu 40 40 Met 45 (2) INFORMATION FOR SEQ ID NO: 476: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 42 amino acids (B) TYPE: amino acid 50 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 476: Pro Arg Leu Ile Arg Gly Arg Val His Arg Cys Val Gly Asn Tyr Asp 55 Gln Lys Lys Asn Ile Phe Gln Cys Val Ser Val Arg Pro Ala Ser Val

Ser Glu Gln Lys Thr Phe Gln Ala Phe Val

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•	35	•			
5	(2) INFORMATION FO	OR SEQ ID NO	: 477:		
	(A) (B)	CE CHARACTER LENGTH: 370 TYPE: amino) amino acid o acid	ls	
10	(xi) SEQUE	TOPOLOGY: 1 NCE DESCRIPI	TION: SEQ ID		
	Gly Val Phe Arg P	ro Cys Val C 5	ys Gly Arg 10	Pro Ala Ser L	eu Thr Cys 15
15	Ser Pro Leu Asp P	ro Glu Val G	Sly Pro Tyr 25	Cys Asp Thr F	ro Thr Met 30
20	Arg Thr Leu Phe A	sn Leu Leu 1	Trp Leu Ala 40	Leu Ala Cys S 45	Ser Pro Val
	His Thr Thr Leu S	Ser Lys Ser i 55	Asp Ala Lys	Lys Ala Ala s 60	Ser Lys Thr
25	Leu Leu Glu Lys 5 65	Ser Gln Phe	Ser Asp Lys	Pro Val Gln 7	Asp Arg Gly 80
30	Leu Val Val Thr	Asp Leu Lys 85	Ala Glu Ser 90	Val Val Leu	Glu His Arg 95
	Ser Tyr Cys Ser 2	Ala Lys Ala	Arg Asp Arg 105	His Phe Ala	Gly Asp Val 110
35	Leu Gly Tyr Val	Thr Pro Trp	Asn Ser His 120	Gly Tyr Asp 125	Val Thr Lys
	Val Phe Gly Ser 130	Lys Phe Thr 135	Gln Ile Ser	Pro Val Trp	Leu Gln Leu
40	Lys Arg Arg Gly 145	Arg Glu Met 150	Phe Glu Val	155 Leu	His Asp Val 160
45	Asp Gln Gly Trp	Met Arg Ala 165	Val Arg Lys	s His Ala Lys O	Gly Leu His 175
	Ile Val Pro Arg 180	Leu Leu Phe	Glu Asp Tr 185	p Thr Tyr Asp	Asp Phe Arg 190
50	Asn Val Leu Asp 195	Ser Glu Asp	Glu Ile Gl 200	u Glu Leu Ser 205	Lys Thr Val
	Val Gln Val Ala 210	Lys Asn Glr 215		p Gly Phe Val 220	Val Glu Val
55	Trp Asn Gln Leu	Leu Ser Glr	n Lys Arg Va	al Gly Leu Ile 235	His Met Leu 240

Thr His Leu Ala Glu Ala Leu His Gln Ala Arg Leu Leu Ala Leu Leu

-	Val	Ile	Pro	Pro 260	Ala	Ile	Thr	Pro	Gly 265	Thr	Asp	Gln	Leu	Gly 270	Met	Phe	е
5	Thr	His	Lys 275	Glu	Phe	Glu	Gln	Leu 280	Ala	Pro	Val	Leu	Asp 285	Gly	Phe	Se	r
	Leu	Met 290	Thr	Tyr	Asp	Tyr	Ser 295	Thr	Ala	His	Gln	Pro 300	Gly	Pro	Asn	Al	a
10	Pro 305	Leu	Ser	Trp	Val	Arg 310	Ala	Cys	Val	Gln	Val 315	Leu	Asp	Pro	Lys	Xa 32	a 0
15				Thr	325					330					335	1	
				9ro 340					345	•				350			
20	Ile	Gln	355	Leu	Lys	Asp	His	360		Arg	Met	: Val	165	Asp	Ser	. Ly	/S
	Pro	Gln 370													٠		
25									470								
	(2)	INE		ATION SEQ													
30					(A) (B) (D)	LENG TYPE TOPO	TH: : am LOGY	39 a ino : li	mino acid near	aci		n. 1	78.			-	
25		_) SE										s Asi	n Th	r P	ro
35		r Cy: 1	s Se	r Pr		u Ası	D PI	O .GI	u va	1		U IJ			1	5	
40	Th	r Me	t Ar	g Th 2		u Ph	e As	n Le	u Le 2		p Le	u Al	a Le	u Al 3	a Cy 0	rs S	er
	Pr	o Va		s Th 5	r Th	r Le	u Se	r									
45	(2	:) IN	FORM	IATI C	n FC	R SE	Q II	NO:	479	:							
50) SE(i) SI	(A) (B) (D)	TYP:	STH: E: a OLOG	54 a mino Y: 1	amino acio inea	o ac: i r		NTO :	179.				
	•	1 <i>7</i> -		al Th										eu G	lu H	is .	Arg
55		1				5					10					72	
	S	er T	AL C	ys Se	er Al 20	la Ly	γs A	la A	rg As	sp A: 25	rg H	is P	ne A	la G	ly A 30	sp	Val
60	L	eu G	ly T	yr V	al T	hr P	ro T	rp A	sn S	er H	is G	ly T	yr A	sp V	al T	hr	Lys

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45 40 35 Val Phe Gly Ser Lys Phe 50 5 (2) INFORMATION FOR SEQ ID NO: 480: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 52 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 480: 15 Arg Glu Met Phe Glu Val Thr Gly Leu His Asp Val Asp Gln Gly Trp 10 Met Arg Ala Val Arg Lys His Ala Lys Gly Leu His Ile Val Pro Arg 20 Leu Leu Phe Glu Asp Trp Thr Tyr Asp Asp Phe Arg Asn Val Leu Asp 25 Ser Glu Asp Glu 50 (2) INFORMATION FOR SEQ ID NO: 481: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 56 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 35 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 481: His Phe Asp Gly Phe Val Val Glu Val Trp Asn Gln Leu Leu Ser Gln 10 5 40 Lys Arg Val Gly Leu Ile His Met Leu Thr His Leu Ala Glu Ala Leu 25 His Gln Ala Arg Leu Leu Ala Leu Leu Val Ile Pro Pro Ala Ile Thr 45 40 35 Pro Gly Thr Asp Gln Leu Gly Met 50 50 (2) INFORMATION FOR SEQ ID NO: 482: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 47 amino acids 55 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 482:

Asp Gly Phe Ser Leu Met Thr Tyr Asp Tyr Ser Thr Ala His Gln Pro

	1	5		10	15
	Gly Pro A	sn Ala Pro 20	Leu Ser Trī	Val Arg Ala C 25	ys Val Gln Val Leu 30
5	Asp Pro I	ys Xaa Lys 35	Trp Arg Th	r Lys Ser Ser 1 O	orp Gly Ser Thr 45
10	(2) INFO	RMATION FOR	SEQ ID NO:	483:	
15		(B) T (D) T	ENGTH: 152 YPE: amino OPOLOGY: 1	amino acids acid	483:
20	1	5		10	Glu Phe Asn Glu Arg 15
	Thr Lys	Asp Ile Lys 20	Glu Gly I	le Pro Leu Pro 25	Thr Lys Ile Leu Val 30
25	Lys Pro	Asp Arg Thr 35	Phe Glu I	le Lys Ile Gly 40	Gln Pro Thr Val Ser 45
30	Tyr Phe 50	Leu Lys Ala	Ala Ala G 55	ly Ile Glu Lys	Gly Ala Arg Gln Thr 60
	Gly Lys 65	Glu Val Ala	a Gly Leu V 70	al Thr Leu Lys 75	His Val Tyr Glu Ile 80
35		8	5	90	Leu Gln Asp Val Pro 95
	Leu Ser	Ser Val Va	l Arg Ser l	lle Ile Gly Ser 105	Ala Arg Ser Leu Gly 110
40	Ile Arg	Val Val Ly 115	s Asp Leu S	Ser Ser Glu Glu 120	Leu Ala Ala Phe Gln 125
٠,	Lys Glu 130		e Phe Leu 1 135	Ala Ala Gln Lys	Glu Ala Asp Leu Ala 140
45	Ala Glr 145	n Glu Glu Al	a Ala Lys 150	Lys	
50	(2) IN	FORMATION FO	OR SEQ ID N	O: 484:	_
		(A)	CE CHARACTI	70 amino acids	
55		(D)	TYPE: amin TOPOLOGY: NCE DESCRI	no acid linear PTION: SEQ ID N	ю: 484:
60	Ala Va	al Tyr Thr T	yr His Glu 5	Lys Lys Lys As	op Thr Ala Ala Ser Gly 15

	Tyr Gly Thr Gln Asn Ile Arg Leu Ser Arg Asp Ala Val Lys Asp Phe 20 25 30
5	Asp Cys Cys Cys Leu Ser Leu Gln Pro Cys His Asp Pro Val Val Thr 35 40 45
	Pro Asp Gly Tyr Leu Tyr Glu Arg Glu Ala Ile Leu Glu Tyr Ile Leu 50 55 60
10	His Gln Lys Lys Glu Ile Ala Arg Gln Met Lys Ala Tyr Glu Lys Gln 65 70 75 80
15	Arg Gly Thr Arg Arg Glu Glu Gln Lys Glu Leu Gln Arg Ala Ala Ser 85 90 95
	Gln Asp His Val Arg Gly Phe Leu Glu Lys Glu Ser Ala Ile Val Ser 100 105 110
20	Arg Pro Leu Asn Pro Phe Thr Ala Lys Ala Leu Ser Gly Thr Ser Pro 115 120 125
25	Asp Asp Val Gln Pro Gly Pro Ser Val Gly Pro Pro Ser Lys Asp Lys 130 135 140
25	Asp Lys Val Leu Pro Ser Phe Trp Ile Pro Ser Leu Thr Pro Glu Ala 145 150 155 160
30	Lys Ala Thr Lys Leu Glu Lys Pro Ser Arg Thr Val Thr Cys Pro Met 165 170 175
	Ser Gly Lys Pro Leu Arg Met Ser Asp Leu Thr Pro Val His Phe Thr 180 185 190
35	Pro Leu Asp Ser Ser Val Asp Arg Val Gly Leu Ile Thr Arg Ser Glu 195 200 205
40	Arg Tyr Val Cys Ala Val Thr Arg Asp Ser Leu Ser Asn Ala Thr Pro 210 215 220
	Cys Ala Val Leu Arg Pro Ser Gly Ala Val Val Thr Leu Glu Cys Val 225 230 235 240
45	Glu Lys Leu Ile Arg Lys Asp Met Val Asp Pro Val Thr Gly Asp Lys 245 250 255
	Leu Thr Asp Arg Asp Ile Ile Val Leu Gln Arg Gly Gly Thr 260 265 270
50	
	(2) INFORMATION FOR SEQ ID NO: 485:
55	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 54 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 485:
60) Tyr Leu Tyr Glu Arg Glu Ala Ile Leu Glu Tyr Ile Leu His Gln Lys

	1	5	1	10	15
_	Lys Glu Ile	Ala Arg Gln M	et Lys Ala Ty 25	yr Glu Lys Glr	a Arg Gly Thr 30
5	Arg Arg Glu 35	Glu Gln Lys G	lu Leu Gln A 40	rg Ala Ala Sei 49	c Gln Asp His
10	Val Arg Gly 50	Phe Leu Glu			
15		TION FOR SEQ I			
		(B) TYPE: (D) TOPOLO	: 64 amino ao amino acid GY: linear		
20	(xi)	SEQUENCE DESC	CRIPTION: SE	Q ID NO: 486:	
	Phe Thr Ala	Lys Ala Leu S 5	Ser Gly Thr S	Ser Pro Asp As 10	p Val Gln Pro 15
25	Gly Pro Ser	Val Gly Pro	Pro Ser Lys 1 25	Asp Lys Asp Ly	rs Val Leu Pro 30
20	Ser Phe Trp		Leu Thr Pro (Glu Ala Lys Al	la Thr Lys Leu 15
30	Glu Lys Pro 50	Ser Arg Thr	Val Thr Cys 55	Pro Met Ser G	ly Lys Pro Leu
35					
40		ATION FOR SEQ			
	(i)	(B) TYPE:	H: 56 amino a amino acid	: acids	
45	(xi	.) SEQUENCE DE	OGY: linear SCRIPTION: SI	EQ ID NO: 487:	:
	Val His Ph 1	e Thr Pro Leu 5	Asp Ser Ser	Val Asp Arg V 10	al Gly Leu Ile 15
50	Thr Arg Se	er Glu Arg Tyr 20	Val Cys Ala 25	Val Thr Arg A	Asp Ser Leu Ser 30
55		nr Pro Cys Ala 35	Val Leu Arg 40	Pro Ser Gly	Ala Val Val Thr 45
J	Leu Glu Cy 50	ys Val Glu Lys	Leu Ile 55		

	(2) INFORMATION FOR SEQ ID NO. 100.
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 567 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 488:
10	Met Asp Thr Ser Glu Asn Arg Pro Glu Asn Asp Val Pro Glu Pro Pro 1 5 10 15
	Met Pro Ile Ala Asp Gln Val Ser Asn Asp Asp Arg Pro Glu Gly Ser 20 25 30
15	Val Glu Asp Glu Glu Lys Lys Glu Ser Ser Leu Pro Lys Ser Phe Lys 35 40 45
	Arg Lys Ile Ser Val Val Ser Ala Thr Lys Gly Val Pro Ala Gly Asn 50 55 60
20	Ser Asp Thr Glu Gly Gln Pro Gly Arg Lys Arg Arg Trp Gly Ala 65 70 75 80
25	Ser Thr Ala Thr Thr Gln Lys Lys Pro Ser Ile Ser Ile Thr Thr Glu 85 90 95
	Ser Leu Lys Ser Leu Ile Pro Asp Ile Lys Pro Leu Ala Gly Gln Glu 100 105 110
30	Ala Val Val Asp Leu His Ala Asp Asp Ser Arg Ile Ser Glu Asp Glu 115 120 125
25	Thr Glu Arg Asn Gly Asp Asp Gly Thr His Asp Lys Gly Leu Lys Ile 130 135 140
35	Cys Arg Thr Val Thr Gln Val Val Pro Ala Glu Gly Gln Glu Asn Gly 145 150 155 160
40	Gln Arg Glu Glu Glu Glu Glu Lys Glu Pro Glu Ala Glu Pro Pro 165 170 175
	Val Pro Pro Gln Val Ser Val Glu Val Ala Leu Pro Pro Pro Ala Glu 180 185 190
45	His Glu Val Lys Lys Val Thr Leu Gly Asp Thr Leu Thr Arg Arg Ser 195 200 205
	Ile Ser Gln Gln Lys Ser Gly Val Ser Ile Thr Ile Asp Asp Pro Val 210 215 220
50	Arg Thr Ala Gln Val Pro Ser Pro Pro Arg Gly Lys Ile Ser Asn Ile 225 230 235 240
55	Val His Ile Ser Asn Leu Val Arg Pro Phe Thr Leu Gly Gln Leu Lys 245 250 255
	Glu Leu Leu Gly Arg Thr Gly Thr Leu Val Glu Glu Ala Phe Trp Il 260 265 270
60	The Tyr Ser The Val Glu Gl

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•	275		280	285	
_	Ala Val Ala Ti 290 .		la Leu His (95	Gly Val Lys Trp 300	Pro Gln Ser
5	Asn Pro Lys Ph 305	ne Leu Cys A 310	la Asp Tyr I	Ala Glu Gln Asp 315	Glu Leu Asp 320
10	Tyr His Arg G	ly Leu Leu V 325	al Asp Arg :	Pro Ser Glu Thr 330	Lys Thr Glu 335
		le Pro Arg F 40	Pro Leu His	Pro Pro Pro Pro	Pro Pro Val 350
15	Gln Pro Pro G 355	ln His Pro A	Arg Ala Glu 360	Gln Arg Glu Gln 365	Glu Arg Ala
20	Val Arg Glu G 370		Glu Arg Glu 375	Arg Glu Met Glu 380	Arg Arg Glu
20	Arg Thr Arg S	er Glu Arg (390	Glu Trp Asp	Arg Asp Lys Val 395	Arg Glu Gly 400
25		405		Arg Arg Arg Lys 410	415
		120	425	Lys Glu Lys Ala	430
30	435		440	Phe Arg Lys Th	5
35	450		455	Asp Ser Gln Il 460	
	465	470		Glu Arg Glu Ly 475	480
40		485		Arg Glu Lys Gl 490	495
٠.		500	505		510
45	515		520	ı Arg Glu Arg As 52	25
50	530		535	g Glu Arg Gly A 540	
50	Arg Arg Asp 545	Thr Lys Arg		g Ser Arg Ser A 555	rg Ser Thr Pro 560
55	Val Arg Asp	Arg Gly Gly 565	/ Arg		

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 489:
5
     Gly Cys Asp Ser Cys Pro Pro His Leu Pro Arg Glu Ala Phe Ala Gln
     Asp Thr Gln Ala Glu Gly Glu Cys Ser Ser Arg Ala Glu Arg Ala Asp
10
      Met Cys Pro Asp Ala Pro Pro Ser Gln Glu Val Pro Glu Gly Pro Gly
                                   40
15
      Ala Ala Pro
           50
20
      (2) INFORMATION FOR SEQ ID NO: 490:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 50 amino acids
25
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 490:
      Pro Gln Leu Pro Ser Cys Gly Arg Pro Trp Pro Gly Thr Ala Ser Val
                                           10
30
                        5
       Phe Gln Ser His Thr Gln Gly Pro Arg Glu Asp Pro Asp Pro Cys Arg
       Ala Gln Gly Ser Ala Gly Thr His Cys Pro Ile Ser Leu Ser Pro Pro
 35
                                   40
       Arg Gln
            50
 40
       (2) INFORMATION FOR SEQ ID NO: 491:
               (i) SEQUENCE CHARACTERISTICS:
 45
                      (A) LENGTH: 42 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 491:
 50
       Pro Gly Phe Arg Gly Pro Ser Gly Ser Leu Gly Cys Ser Phe Pro
       Arg Ser Leu Gly Arg Val Leu Pro Pro Gly Cys Gln Arg Pro Gly Ala
 55
       His Ala Asp Ser Ser Pro Pro Pro Thr Pro
```

	(2) INFORMATION FOR SEQ ID No. 432.									
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 84 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear									
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 492:									
10	Glu Asp Leu Lys Lys Pro Asp Pro Ala Ser Leu Arg Ala Ala Ser Cys 1 5 10 15									
1.5	Gly Glu Gly Lys Lys Arg Lys Ala Cys Lys Asn Cys Thr Cys Gly Leu 20 25 30 .									
15	Ala Glu Glu Leu Glu Lys Glu Lys Ser Arg Glu Gln Met Ser Ser Gln 35 40 45									
20	Pro Lys Ser Ala Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe Arg Cys 50 55 60									
	Ala Ser Cys Pro Tyr Leu Gly Met Pro Ala Phe Lys Pro Gly Glu Lys 65 70 75 80									
25	Val Leu Leu Ser									
30	(2) INFORMATION FOR SEQ ID NO: 493:									
	(i) SEQUENCE CHARACTERISTICS:									
	(A) LENGTH: 90 amino acids (B) TYPE: amino acid									
35	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 493:									
40	Glu Asp Leu Lys Lys Pro Asp Pro Ala Ser Leu Arg Ala Ala Ser Cys 1 5 10 15									
40	Gly Glu Gly Lys Lys Arg Lys Ala Cys Lys Asn Cys Thr Cys Gly Leu 20 25 30									
45	Ala Glu Glu Leu Glu Lys Glu Lys Ser Arg Glu Gln Met Ser Ser Gln 35 40 45									
	Pro Lys Ser Ala Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe Arg Cys 50 55 60									
50	Ala Ser Cys Pro Tyr Leu Gly Met Pro Ala Phe Lys Pro Gly Glu Lys 65 70 75 86									
55	Val Leu Leu Ser Asp Ser Asn Leu His Asp 85 90									
	(2) INFORMATION FOR SEQ ID NO: 494:									
60	(i) SEQUENCE CHARACTERISTICS:									

ASD

```
(A) LENGTH: 34 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 494:
5
     Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe Arg Cys Ala Ser Cys Pro
     Tyr Leu Gly Met Pro Ala Phe Lys Pro Gly Glu Lys Val Leu Leu Ser
10
      Asp Ser
15
      (2) INFORMATION FOR SEQ ID NO: 495:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 25 amino acids
20
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 495:
       Ser Cys Gly Glu Gly Lys Lys Arg Lys Ala Cys Lys Asn Cys Thr Cys
 25
                                            10
       Gly Leu Ala Glu Glu Leu Glu Lys Glu
                    20
 30
       (2) INFORMATION FOR SEQ ID NO: 496:
               (i) SEQUENCE CHARACTERISTICS:
 35
                     (A) LENGTH: 21 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 496:
  40
        Ser Gln Pro Lys Ser Ala Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe
                                             10
        Arg Cys Ala Ser Cys
  45
                     20
         (2) INFORMATION FOR SEQ ID NO: 497:
   50
                (i) SEQUENCE CHARACTERISTICS:
                       (A) LENGTH: 17 amino acids
                        (B) TYPE: amino acid
                        (D) TOPOLOGY: linear
                (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 497:
   55
         Arg Glu Ala Gly Gln Asn Ser Glu Arg Gln Tyr Val Ser Leu Ser Arg
                                              10
   60
```

5	(2) INFORMATION FOR SEQ ID NO: 498:										
10	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 90 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 498: 										
	Glu Ser Ser Gly Gln Ala Arg Thr Leu Ala Asp Pro Gly Pro Gly Trp 1 5 10 15										
15	Pro Arg Gln Gln Gly Met Cys Phe Gly Ser Leu Thr Gly Leu Ser Thr 20 25 30										
20	Thr Pro His Gly Phe Leu Thr Val Ser Ala Glu Ala Asp Pro Arg Leu 35 40 45										
	Ile Glu Ser Leu Ser Gln Met Leu Ser Met Gly Phe Ser Asp Glu Gly 50 55 60										
25	Gly Trp Leu Thr Arg Leu Leu Gln Thr Lys Asn Tyr Asp Ile Gly Ala 65 70 75 80										
	Ala Leu Asp Thr Ile Gln Tyr Ser Lys His 85 90										
30											
	(2) INFORMATION FOR SEQ ID NO: 499:										
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 159 amino acids (B) TYPE: amino acid										
40	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 499:										
40	Gln Glu Gly Ser Glu Pro Val Leu Leu Glu Gly Glu Cys Leu Val Val 1 5 10 15										
45	Cys Glu Pro Gly Arg Ala Ala Ala Gly Gly Pro Gly Gly Ala Ala Leu 20 25 30										
	Gly Glu Ala Pro Pro Gly Arg Val Ala Phe Xaa Ala Val Arg Ser His 35 40 45										
50	His His Glu Pro Ala Gly Glu Thr Gly Asn Gly Thr Ser Gly Ala Ile 50 55 60										
55	Tyr Phe Asp Gln Val Leu Val Asn Glu Gly Gly Fhe Asp Arg Ala 65 70 75 80										
55	Ser Gly Ser Phe Val Ala Pro Val Arg Gly Val Tyr Ser Phe Arg Phe 85 90 95										
60	His Val Val Lys Val Tyr Asn Arg Gln Thr Val Gln Val Ser Leu Met										

	Leu Asn Thr Trp Pro Val Ile Ser Ala Phe Ala Asn Asp Pro Asp Val 115 120 125
5	Thr Arg Glu Ala Ala Thr Ser Ser Val Leu Leu Pro Leu Asp Pro Gly 130 135 140
10	Asp Arg Val Ser Leu Arg Leu Arg Arg Gly Xaa Ser Thr Gly Trp 145 150 155
	(2) INFORMATION FOR SEQ ID NO: 500:
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 32 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 500: Pro Arg Ser Arg Pro Ala Leu Arg Pro Gly Arg Gln Arg Pro Pro Ser 1 5 10 15
25	His Ser Ala Thr Ser Gly Val Leu Arg Pro Arg Lys Lys Pro Asp Pro 20 25 30
30	
	(2) INFORMATION FOR SEQ ID NO: 501: (i) SEQUENCE CHARACTERISTICS:
35	(A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 501:
40	Met Thr Leu Ile Thr Pro Ser Xaa Lys Leu Thr Phe Xaa Lys Gly Asn 1 5 10 15
45	Lys Ser Trp Ser Ser Arg Ala Cys Ser Ser Thr Leu Val Asp Pro 20 25 30
	(2) INFORMATION FOR SEQ ID NO: 502:
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 51 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 502:
	Gly His Pro Ser Pro Ala Leu Ser Ile Ala Pro Ser Asp Gly Ser Gin 1 5 10 15
60	Leu Pro Cys Asp Glu Val Pro Tyr Gly Glu Ala His Val Thr Arg Tyr 20 25 30

	35 40 45	
5	Ser Ser Leu 50	
10	(2) INFORMATION FOR SEQ ID NO: 503:	•
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 503:	
20	GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC	60
	GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	120
26	CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	180
25	TCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	240
	AAGTATTAAA AGTAGCTTTG TAA	263
30		
	(2) INFORMATION FOR SEQ ID NO: 504:	
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double	
35 40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	6
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504:	6 12
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	12
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	12
40 45 50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC TCCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	12 18 24
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC TCCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	12 18 24
40 45 50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 263 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504: GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC TCCCCGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA AAGTATTAAA AGTAGCTTTG TAA	12 18 24

	(C) STRANDEDNESS: double (D) TOPOLOGY: linear										
_	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 505:										
5	GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC										
	GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	120									
10	CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	180									
	TCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	240									
15	AAGTATTAAA AGTAGCTTTG TAA	263									
20	(2) INFORMATION FOR SEQ ID NO: 506: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 160 base pairs										
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear										
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 506:										
30	TOGCTCACTG TCTTACAATC ACTGCTGTGG AATCATGATA CCACTTTTAG CTCTTTGCAT										
	CITCCTTCAG TGTATTTTTG TTTTTCAAGA GGAAGTAGAT TTTAACTGGA CAACTTTGAG	120 160									
35	TACTGACATC ATTGATAAAT AAACTGGCTT GTGGTTTCAA	160									
40	(2) INFORMATION FOR SEQ ID NO: 507: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 292 amino acids (B) TYPE: amino acid										
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 507: Leu Asp Glu Leu Met Ala His Leu Thr Glu Met Gln Ala Lys Val Ala 1 5 10 15										
50											
	Lys Ala Ser Leu Asp Ser Met Leu Gly Gly Leu Glu Gln Glu Leu Gln 35 40 45										
55	50 55 60										
60	Lys Pro Ile Ala Gly Lys Val Ile His Ala Leu Gly Gln Ser Trp His 65 70 75 80										

	Pro Glu His Phe Val Cys Thr His Cys Lys Glu Glu Ile Gly Ser Ser 85 90 95
5	Pro Phe Phe Glu Arg Ser Gly Leu Xaa Tyr Cys Pro Asn Asp Tyr His 100 105 110
	Gln Leu Phe Ser Pro Arg Cys Ala Tyr Cys Ala Ala Pro Ile Leu Asp 115 120 125
10	Lys Val Leu Thr Ala Met Asn Gln Thr Trp His Pro Glu His Phe Phe 130 135 140
1.5	Cys Ser His Cys Gly Glu Val Phe Gly Ala Glu Gly Phe His Glu Lys 145 150 155 160
15	Asp Lys Lys Pro Tyr Cys Arg Lys Asp Phe Leu Ala Met Phe Ser Pro 165 170 175
20	Lys Cys Gly Gly Cys Asn Arg Pro Val Leu Glu Asn Tyr Leu Ser Ala 180 185 190
	Met Asp Thr Val Trp His Pro Glu Cys Phe Val Cys Gly Asp Cys Phe 195 200 205
25	Thr Ser Phe Ser Thr Gly Ser Phe Phe Glu Leu Asp Gly Arg Pro Phe 210 215 220
20	Cys Glu Leu His Tyr His His Arg Arg Gly Thr Leu Cys His Gly Cys 225 230 235 240
30	Gly Gln Pro Ile Thr Gly Arg Cys Ile Ser Ala Met Gly Tyr Lys Phe 245 250 255
35	His Pro Glu His Phe Val Cys Ala Phe Cys Leu Thr Gln Leu Ser Lys 260 265 270
	Gly Ile Phe Arg Glu Gln Asn Asp Lys Thr Tyr Cys Gln Pro Cys Phe 275 280 285
40	Asn Lys Leu Phe 290
	TO T
45	(2) INFORMATION FOR SEQ ID NO: 508: (i) SEQUENCE CHARACTERISTICS:
50	(A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 508: Lys Ala Ser Leu Asp Ser Met Leu Gly Gly Leu Glu Gln Glu Leu Gln
55	1 5 10 13 Asp Leu Gly Ile Ala Thr Val Pro Lys Gly His Cys Ala Ser Cys Glr
	20 25 30 Lys Pro Ile Ala Gly Lys Val Ile His Ala Leu
60	35 40

_	(2) INFORMATION FOR SEQ ID NO: 509:
5	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 50 amino acids
	(B) TYPE: amino acid
	(D) TOPOLOGY: linear
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 509:
	Cys Pro Asn Asp Tyr His Gln Leu Phe Ser Pro Arg Cys Ala Tyr Cys 1 5 10 15
15	Ala Ala Pro Ile Leu Asp Lys Val Leu Thr Ala Met Asn Gln Thr Trp 20 25 30
00	His Pro Glu His Phe Phe Cys Ser His Cys Gly Glu Val Phe Gly Ala 35 40 45
20	Glu Gly 50
25	(2) INFORMATION FOR SEQ ID NO: 510:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 67 amino acids
30	(B) TYPE: amino acid
30	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 510:
	The Dear Die Loui Ala Met Phe Ser Pro
35	Asp Lys Lys Pro Tyr Cys Arg Lys Asp Phe Leu Ala Met Phe Ser Pro
22	*
	Lys Cys Gly Gly Cys Asn Arg Pro Val Leu Glu Asn Tyr Leu Ser Ala 20 25 30
40	Met Asp Thr Val Trp His Pro Glu Cys Phe Val Cys Gly Asp Cys Phe
40	35 40 45
	Thr Ser Phe Ser Thr Gly Ser Phe Phe Glu Leu Asp Gly Arg Pro Phe
	50 55 60
45	Cys Glu Leu
	65
	•
	·
50	(2) INFORMATION FOR SEQ ID NO: 511:
-	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 46 amino acids
55	(B) TYPE: amino acid
	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 511:
	Cys Gly Gln Pro Ile Thr Gly Arg Cys Ile Ser Ala Met Gly Tyr Lys
60	

	Phe His Pro Glu His Phe Val Cys Ala Phe Cys Leu Thr Gli Leu Ser 20 25 30
5	Lys Gly Ile Phe Arg Glu Gln Asn Asp Lys Thr Tyr Cys Gln 35 40 45
10	(2) INFORMATION FOR SEQ ID NO: 512:
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 452 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 512:
20	Met Gly Ser Ser Gln Ser Val Glu Ile Pro Gly Gly Gly Thr Glu Gly 1 5 10 15 Tyr His Val Leu Arg Val Gln Glu Asn Ser Pro Gly His Arg Ala Gly
	20 25 30
25	Leu Glu Pro Phe Phe Asp Phe Ile Val Ser Ile Asn Gly Ser Arg Leu 35 40 45
	Asn Lys Asp Asn Asp Thr Leu Lys Asp Leu Leu Lys Xaa Asn Val Glu 50 55 60
30	Lys Pro Val Lys Met Leu Ile Tyr Ser Ser Lys Thr Leu Glu Leu Arg 65 70 75 80
35	Glu Thr Ser Val Thr Pro Ser Asn Leu Trp Gly Gly Gln Gly Leu Leu 85 90 95
33	Gly Val Ser Ile Arg Phe Cys Ser Phe Asp Gly Ala Asn Glu Asn Val 100 105 110
40	Trp His Val Leu Glu Val Glu Ser Asn Ser Pro Ala Ala Leu Ala Gly 115 120 125
	Leu Arg Pro His Ser Asp Tyr Ile Ile Gly Ala Asp Thr Val Met Asn 130 135 140
45	Glu Ser Glu Asp Leu Phe Ser Leu Ile Glu Thr His Glu Ala Lys Pro 145 150 155 160
50	Leu Lys Leu Tyr Val Tyr Asn Thr Asp Thr Asp Asn Cys Arg Glu Val 165 170 175
30	Ile Ile Thr Pro Asn Ser Ala Trp Gly Gly Glu Gly Ser Leu Gly Cys 180 185 190
55	Gly Ile Gly Tyr Gly Tyr Leu His Arg Ile Pro Thr Arg Pro Phe Glu 195 200 205
	Glu Gly Lys Lys Ile Ser Leu Pro Gly Gln Met Ala Gly Thr Pro Ile 210 215 220 .
60	Thr Pro Leu Lys Asp Gly Phe Thr Glu Val Gln Leu Ser Ser Val Asn

	225					230					235					240
_	Pro	Pro	Ser	Leu	Ser 245	Pro	Pro	Gly	Thr	Thr 250	Gly	Ile	Glu	Gln	Ser 255	Leu
5	Thr	Gly	Leu	Ser 260	Ile	Ser	Ser	Thr	Pro 265	Pro	Ala	Val	Ser	Ser 270	Val	Leu
10	Ser	Thr	Gly 275	Val	Pro	Thr	Val	Pro 280	Leu	Leu	Pro	Pro	Gln 285	Val	Asn	Gln
	Ser	Leu 290	Thr	Ser	Val	Pro	Pro 295	Met	Asn	Pro	Ala	Thr 300	Thr	Leu	Pro	Gly
15	Leu 305	Met	Pro	Leu	Pro	Ala 310	Gly	Leu	Pro	Asn	Leu 315	Pro	Asn	Leu	Asn	Leu 320
20	Asn	Leu	Pro	Ala	Pro 325	His	Ile	Met	Pro	330		Gly	Leu	Pro	Glu 335	Leu
20	Val	Asn	Pro	Gly 340		Pro	Pro	Leu	Pro 345		Met	Pro	Pro	Arg 350	Asn	Leu
25	Pro	Gly	Ile 355		Pro	Leu	Pro	Leu 360		Ser	Glu	Phe	Leu 365	Pro	Ser	Phe
	Pro	Leu 370		Pro	Glu	Ser	Ser 375		Ala	Ala	Ser	Ser 380	-	Glu	Leu	Leu
30	Ser 385		Leu	Pro	Pro	Thr 390		Asn	Ala	Pro	Ser 395		Pro	Ala	Thr	Thr 400
35	Thr	Ala	Lys	Ala	Asp 405		Ala	Ser	Ser	410		Val	Asp	Val	Thr 415	Pro
	Pro	Thr	Ala	Lys 420		Pro	Thr	Thr	Val 425		Asp	Arg	Val	Gly 430		Ser
40	Thr	Pro	Val 435		Glu	Lys	Pro	Val 440		: Ala	a Ala	. Val	445		Asn	Ala
-	Ser	Glu 450		Pro	•											
45	(2)	TNF	ORMZ	ኒ ጥፐ ርነ እ	I FOE	SPO) TD	NO:	513:							
	(-,							TERIS								
50			(xi		(B) '	TYPE TOPO	: am	109 a ino a : li :PTIC	acid near			D: 5	13:			
55		r Vai	l Glı	ı Ile	_	Gl ₃	/ Gl	y Gly	Th i	r Ġlı		у Тул	r His	Va.	Leu 15	ı Arg
60	Va]	l Gl	n Glu	ı Ası 20		r Pro	Gl ₃	y His	Arq 2		a Gly	/ Le	ı Glı	ı Pro		e Phe

	Asp	Phe	Ile 35	Val	Ser	Ile	Asn	Gly 40	Ser	Arg	Leu	Asn	Lys 45	Asp	Asn	Asp
5	Thr	Leu 50	Lys	Asp	Leu	Leu	Lys 55	Xaa	Asn	Val	Glu	Lys 60	Pro	Val	Lys	Met
	Leu 65	Ile	Tyr	Ser	Ser	Lys 70	Thr	Leu	Glu	Leu	Arg 75	Glu	Thr	Ser	Val	Thr 80
10	Pro	Ser	Asn	Leu	Trp 85	Gly	Gly	Gln	Gly	Leu 90	Leu	Gly	Val	Ser	Ile 95	Arg
15	Phe	Cys	Ser	Phe 100	Asp	Gly	Ala	Asn	Glu 105	Asn	Val	Trp	His			
	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: !	514:							
20			(i)	(ENGT YPE :	H: 1 ami	45 a no a	mino .cid	: aci	ds					
25	-1			SEQ											C	3
	1		ASN	Ser	Pro 5	Ala	Ala	Leu	Ala	10	Leu	Arg	PIO	HIS	15	Asp
30	Tyr	Ile	Ile	Gly 20	Ala	Asp	Thr	Val	Met 25	Asn	Glu	Ser	Glu	Asp 30	Leu	Phe
	Ser	Leu	Ile 35		Thr	His	Glu	Ala 40		Pro	Leu	Lys	Leu 45		Val	Tyr
35	Asn	Thr 50	_	Thr	Asp	Asn	Cys 55		Glu	Val	Ile	Ile 60	Thr	Pro	Asn	Ser
40	Ala 65	_	Gly	Gly	Glu	Gly 70		Leu	Gly	Cys	Gly 75		Gly	Tyr	Gly	Tyr 80
	Leu	His	Arg	Ile	Pro 85		Arg	Pro	Phe	Glu 90		Gly	Lys	Lys	Ile 95	Ser
45	Lev	Pro	Gly	Gln 100		Ala	Gly	Thr	Pro 105		Thr	Pro	Leu	Lys 110	_	Gly
	Phe	. Thr	Glu 115		Gln	Leu	Ser	Ser 120		. Asn	Pro	Pro	Ser 125		Ser	Pro
50	Pro	Gly 130		Thr	Gly	lle	Glu 135		Sex	Leu	Thr	Gly 140		Ser	Ile	e Ser
55	Ser 145															
	(2)	IN	PORM	ATION	FOF	SEC	ID.	NO:	515:							

(i) SEQUENCE CHARACTERISTICS:

	(A) LENGTH: 145 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 515:															
5			(xi)							EQ II	ОИС	515	i:			
	Glu 1	Ser	Asn	Ser	Pro 5	Ala	Ala	Leu	Ala	Gly 10	Leu	Arg	Pro	His	Ser 15	qzA
10	Tyr	Ile	Ile	Gly 20	Ala	Asp	Thr	Val	Met 25	Asn	Glu	Ser	Glu	Asp 30	Leu	Phe
	Ser	Leu	Ile 35	Glu	Thr	His	Glu	Ala 40	Lys	Pro	Leu	Lys	Leu 45	Tyr	Val	Tyr
15	Asn	Thr 50	Asp	Thr	Asp	Asn	Cys 55	Arg	Glu	Val	Ile	Ile 60	Thr	Pro	Asn	Ser
20	Ala 65	Trp	Gly	Gly	Glu	Gly 70	Ser	Leu	Gly	Суз	Gly 75	Ile	Gly	Tyr	Gly	Tyr 80
	Leu	His	Arg	Ile	Pro 85	Thr	Arg	Pro	Phe	Glu 90	Glu	Gly	Lys	Lys	Ile 95	Ser
25	Leu	Pro	Gly	Gln 100	Met	Ala	Gly	Thr	Pro 105	Ile	Thr	Pro	Leu	Lys 110	Asp	Gly
	Phe	Thr	Glu 115	Val	Gln	Leu	Ser	Ser 120	Val	Asn	Pro	Pro	Ser 125	Leu	Ser	Pro
30	Pro	Gly 130		Thr	Gly	Ile	Glu 135		Ser	Leu	Thr	Gly 140	Leu	Ser	Ile	Ser
35	Ser 145															
	(2)	INF	ORMA	TION	FOR	SEQ	IĎ	NO:	516:							
40			(i)		(A) I	ENG	TH: 3	TERIS 151 a ino a	mino		ids					
			(xi)	SEC				: lir CPTIC		EQ 1	D NC	: 51	6:			
45	Arg		Pro	Thr	Arg		Phe	e Glu	Glu	Gly 10		Lys	Ile	Ser	Leu 15	
50	Gly	Gln	Met	Ala 20	_	Thr	Pro	Ile	Thr 25		Leu	Lys	Asp	Gly 30	Phe	Thr
	Glu	Va]	. Glr 35		Ser	Ser	(Va)	L Asn		Pro	Ser	Leu	Ser 45		Pro	Gly
55	Thr	Thr 50	-	/ Ile	: Glu	Glr	Ser 55		Thr	Gly	r Leu	Sex 60		. Ser	Ser	Thr
60	Pro 65		Ala	a Val	. Sei	: Se:	_	l Leu	Ser	Thi	Gl ₃		. Pro	Thr	Val	Pro 80

	Leu Leu Pro Pro Gln Val Asn Gln Ser Leu Thr Ser Val Pro Pro Met 85 90 95
5	Asn Pro Ala Thr Thr Leu Pro Gly Leu Met Pro Leu Pro Ala Gly Leu 100 105 110
	Pro Asn Leu Pro Asn Leu Asn Leu Asn Leu Pro Ala Pro His Ile Met 115 120 125
10	Pro Gly Val Gly Leu Pro Glu Leu Val Asn Pro Gly Leu Pro Pro Leu 130 135 140
15	Pro Ser Met Pro Pro Arg Asn 145 150
	(2) INFORMATION FOR SEQ ID NO: 517:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 109 amino acids (B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 517:
25	Pro Gly Leu Pro Pro Leu Pro Ser Met Pro Pro Arg Asn Leu Pro Gly 1 5 10 15
30	Ile Ala Pro Leu Pro Leu Pro Ser Glu Phe Leu Pro Ser Phe Pro Leu 20 25 30
	Val Pro Glu Ser Ser Ser Ala Ala Ser Ser Gly Glu Leu Leu Ser Ser 35 40 45
35	Leu Pro Pro Thr Ser Asn Ala Pro Ser Asp Pro Ala Thr Thr Thr Ala 50 55 60
40	Lys Ala Asp Ala Ala Ser Ser Leu Thr Val Asp Val Thr Pro Pro Thr 65 70 75 80
	Ala Lys Ala Pro Thr Thr Val Glu Asp Arg Val Gly Asp Ser Thr Pro 85 90 95
45	Val Ser Glu Lys Pro Val Ser Ala Ala Val Asp Ala Asn 100 105
50	(2) INFORMATION FOR SEQ ID NO: 518:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 93 amino acids
<i>e</i>	(B) TYPE: amino acid (D) TOPOLOGY: linear
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 518:
	Ile Tyr Lys Val Phe Arg His Thr Ala Gly Leu Lys Pro Glu Val Ser 1 5 10 15
60	Cys Phe Glu Asn Ile Arg Ser Cys Ala Arg Xaa Xaa Xaa Xaa Xaa

629

20 25 30 Xaa Xaa Xaa Xaa Xaa Trp Ile Phe Gly Val Leu His Val Val His 40 5 Ala Ser Val Val Thr Ala Tyr Leu Phe Thr Val Ser Asn Ala Phe Gln 55 Gly Met Phe Ile Phe Leu Phe Leu Cys Val Leu Ser Arg Lys Ile Gln 10 Glu Glu Tyr Tyr Arg Leu Phe Lys Asn Val Pro Cys Cys 85 15 (2) INFORMATION FOR SEQ ID NO: 519: (i) SEQUENCE CHARACTERISTICS: 20 (A) LENGTH: 55 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 519: Trp Ile Phe Gly Val Leu His Val Val His Ala Ser Val Val Thr Ala 25 Tyr Leu Phe Thr Val Ser Asn Ala Phe Gln Gly Met Phe Ile Phe Leu 30 Phe Leu Cys Val Leu Ser Arg Lys Ile Gln Glu Glu Tyr Tyr Arg Leu Phe Lys Asn Val Pro Cys Cys 35 50 (2) INFORMATION FOR SEQ ID NO: 520: 40 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 50 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 520: Ala Leu Thr Arg Ile Pro Pro Gly Asp Trp Val Ile Asn Val Thr Ala 50 Val Ser Phe Ala Gly Lys Thr Thr Ala Arg Phe Phe Xaa His Ser Ser 25 Pro Pro Ser Leu Gly Asp Gln Ala Arg Thr Asp Pro Gly His Gln Arg 40 55 Arg Asp 50

•	(2) INFORMATION FOR SEQ ID NO: 521:	
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 521:	
10	Leu Gln Glu Val Asn Ile Thr Leu Pro Glu Asn Ser Val Trp Tyr Glu 1 5 10 15	
	Arg Tyr Lys Phe Asp Ile Pro Val Phe His Leu 20 25	
15		
20	(2) INFORMATION FOR SEQ ID NO: 522: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 110 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 522:	
25	Met Gln Gly Ser Gly Ser Gln Phe Arg Ala Cys Leu Leu Cys Leu Cys 1 5 10 15	
30	Phe Ser Cys Pro Cys Ser Pro Gly Gly Pro Arg Trp Asn Ser Arg Gln 20 25 30	
	Gly Gly Arg Arg Phe Pro Lys Thr Cys Arg Ala Ile Ser Gln Asn Leu 35 40 45	
35	Val Phe Lys Tyr Lys Thr Phe Cys Pro Val Arg Tyr Met Gln Pro His 50 55 60	
	Arg Ser Ser Leu Cys Leu His Phe Thr Ser Tyr Val Phe Ile Leu Ser 65 70 75 80	
40	Thr Trp Gly Ser Leu Arg Thr Tyr Ser Thr Asp Leu Lys Lys Lys Lys 85 90 95	
45	Lys Asn Ser Arg Gly Gly Pro Val Pro Ile Arg Pro Lys Ser 100 105 110	
	(2) INFORMATION FOR SEQ ID NO: 523:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LEWGTH: 99 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 523:	
60	TAGCATGTAG CCAGTCGAAT AACNIATAAG GACAAAGTGG AGTCCACGCG TGCGGCCGTC	60
	mini ami ama armananana amani naram nanari 1010	F14

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(2) INFORMATION FOR SEQ ID NO: 524:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
                    (B) TYPE: amino acid
10
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 524:
     Met Gln Gly Ser Gly Ser Gln Phe Arg Ala Cys Leu Cys Leu Cys
15
      Phe Ser Cys Pro Cys Ser Pro Gly Gly Pro Arg Trp Asn Ser Arg Gln
                                      25
      Gly Gly Arg Arg Phe Pro Lys Thr Cys Arg Ala Ile Ser Gln Asn Leu
20
      Val Phe Lys
           50
25
      (2) INFORMATION FOR SEQ ID NO: 525:
              (i) SEQUENCE CHARACTERISTICS:
30
                     (A) LENGTH: 54 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 525:
      Pro Val Arg Tyr Met Gln Pro His Arg Ser Ser Leu Cys Leu His Phe
35
                                           10
                        5
      Thr Ser Tyr Val Phe Ile Leu Ser Thr Trp Gly Ser Leu Arg Thr Tyr
                                      25
40
      Ser Thr Asp Leu Lys Lys Lys Lys Asn Ser Arg Gly Gly Pro Val
                                    40
       Pro Ile Arg Pro Lys Ser
 45
           50
       (2) INFORMATION FOR SEQ ID NO: 526:
 50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 38 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
 55
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 526:
       Gly Glu Glu Gln Arg Asp Cys Ser Leu Gly Trp Arg Gly Val Gly Met
                                                                15 .
                                            10
       Arg Ala Thr His Cys Gln Ala Ala Arg Met Phe Val Leu Phe Ser Leu
 60
```

632

30 25 20 Pro Lys Tyr Ala Gly Leu 35 5 (2) INFORMATION FOR SEQ ID NO: 527: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 161 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 527: 15 Met Pro Arg Lys Thr Ser Lys Cys Arg Gln Leu Leu Cys Ser Gly Ala Ser Arg Asn Ala Asp Thr Ala Ala Arg Gln Ser Thr Cys Ser Ser His 20 25 Arg Pro Pro Gly Lys Ile Pro Ser Leu Gly Pro Arg Arg Xaa Pro Gly Cys Xaa Ser Val Pro Ser Ser Arg Gly Glu Gln Ser Thr Gly Ser Pro 25 Ala Ala Pro Arg Cys Gly Arg Arg Asp Ala His Arg Gly Leu Pro Gly 30 Gly Ala Ala Met Thr Pro Gly Asp Thr Trp Ala Ser Phe Asn Pro Arg Ala Gly His Ser Lys Ser Gln Gly Glu Gly Gln Glu Ser Ser Gly Ala 105 35 Ser Arg Gln Asp Arg His Pro Val Ser His Trp Val Glu Arg Gln Arg Glu Ala Trp Gly Ala Pro Arg Ser Ser Ser Ala Gly Gly Val Lys Val 40 135 Ala Ala Thr Thr Glu Arg Glu Pro Glu Phe Lys Ile Lys Thr Gly Lys 150 45 Ala 50 (2) INFORMATION FOR SEQ ID NO: 528: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 88 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 528:

5

60

Cys Ser Gly Ala Ser Arg Asn Ala Asp Thr Ala Ala Arg Gln Ser Thr

•	Cys Ser	Ser H	is Ar	g Pro	Pro	Gly	Lys 25	Ile	Pro	Ser	Leu	Gly:	Pro .	Arg
5	Arg Xaa	Pro G	Sly Cy	s Xaa	Ser	Val 40	Pro	Ser	Ser	Arg	Gly 45	Glu	Gln	Ser
10	Thr Gly 50	Ser I	Pro Al	a Ala	Pro 55	Arg	Cys	Gly	Arg	Arg 60	Asp	Ala	His	Arg
10	Gly Leu 65	Pro (Gly Gl	y Ala 70		Met	Thr	Pro	Gly 75	Asp	Thr	Trp	Ala	Ser 80
15	Phe Asn	Pro i		a Gly 5	His	Ser								
20	(2) INF		EQUENC	E CHA	RACI	TERIS	TICS		ls					
25		(xi)		TYPE TOPO:	LOGY	: lir	near	EQ I	D NO): 52	9 :			
	Gln Gly 1	Glu	Gly G	ln Glu 5	ı Ser	Ser	Gly	Ala 10		Arg	Gln	Asp	Arg 15	His
30	Pro Val	Ser	His T	op Va	l Glu	ı Arg	Gln 25		Glu	Ala	Trp	Gly 30	Ala	Pro
35	Arg Ser	Ser 35	Ser A	la Gl	y G13	/ Val		: Val	Ala	Ala	Thr 45		Glu	Arg
	Glu Pro		Phe L	ys Il	e Ly: 5!		c Gly	, Lys	: Ala	1				
40	(2) IN	FORMA:	rion f	OR SE	Q ID	NO:	530	:						
45		•	(B)	TYPE TOP	FTH: E: an OLOGY	235 ino : li	amin acid near	o ac		o: 5	30:			
50	Met Se	r Pro	Arg T	yr Pr 5	o Gl	y Gl	y Pr	o Ar		o Pr	o Le	u Arg	Ile 19	
	Asn Gl	n Ala	Leu 0	ly Gl	y Va	l Pr	o G1 2		r Gl	n Pr	o Le	u Leu 30		Ser
55	Gly Me	t Asp 35		hr Ar	g G1		n Gl	y Hi	s Pr	o As		t Gly 5	/ Gly	y Pro
60	Met Gl 5	n Arg O	Met 7	Thr Pi		ro Ar i5	g Gl	y Me	t Va		o Le	u Gly	y Pro	o Gln

-	Asn Tyr Gly Gly Ala Met Arg Pro Pro Leu Asn Ala Leu Gly Gly Pro 65 70 75 80
5	Gly Met Pro Gly Met Asn Met Gly Pro Gly Gly Gly Arg Pro Trp Pro 85 90 95
	Asn Pro Thr Asn Ala Asn Ser Ile Pro Tyr Ser Ser Ala Ser Pro Gly 100 105 110
10	Asn Tyr Val Gly Pro Pro Gly Gly Gly Gly Pro Pro Gly Thr Pro Ile 115 120 125
15	Met Pro Ser Pro Ala Asp Ser Thr Asn Ser Gly Asp Asn Met Tyr Thr 130 135 140
	Leu Met Asn Ala Val Pro Pro Gly Pro Asn Arg Pro Asn Phe Pro Met 145 150 155 160
20	Gly Pro Gly Ser Asp Gly Pro Met Gly Gly Leu Gly Gly Met Glu Ser 165 170 175
	His His Met Asn Gly Ser Leu Gly Ser Gly Asp Met Asp Ser Ile Ser 180 185 190
25	Lys Asn Ser Pro Asn Asn Met Ser Leu Ser Asn Gln Pro Gly Thr Pro 195 200 205
30	Arg Asp Asp Gly Glu Met Gly Gly Asn Phe Leu Asn Pro Phe Gln Ser 210 215 220
	Glu Ser Tyr Ser Pro Ser Met Thr Met Ser Val 225 230 235
35	(2) INFORMATION FOR SEQ ID NO: 531:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 114 amino acids
40	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 531:
45	Met Ser Pro Arg Tyr Pro Gly Gly Pro Arg Pro Pro Leu Arg Ile Pro
43	Asn Gln Ala Leu Gly Gly Val Pro Gly Ser Gln Pro Leu Leu Pro Ser
50	Gly Met Asp Pro Thr Arg Gln Gln Gly His Pro Asn Met Gly Gly Pro 35 40 45
	Met Gln Arg Met Thr Pro Pro Arg Gly Met Val Pro Leu Gly Pro Gln 50 55 60
55	
60	Gly Met Pro Gly Met Asn Met Gly Pro Gly Gly Gly Arg Pro Trp Pro

	Asn Pro Thr Asn Ala Asn Ser Ile Pro Tyr Ser Ser Ala Ser Pro Gly 100 105 110
5	Asn Tyr
10	(2) INFORMATION FOR SEQ ID NO: 532:
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 81 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 532:
20	Leu Asn Ala Leu Gly Gly Pro Gly Met Pro Gly Met Asn Met Gly Pro 1 5 10 15
	Gly Gly Gly Arg Pro Trp Pro Asn Pro Thr Asn Ala Asn Ser Ile Pro 20 25 30
25	Tyr Ser Ser Ala Ser Pro Gly Asn Tyr Val Gly Pro Pro Gly Gly Gly 35 40 45
	Gly Pro Pro Gly Thr Pro Ile Met Pro Ser Pro Ala Asp Ser Thr Asn 50 55 60
30	Ser Gly Asp Asn Met Tyr Thr Leu Met Asn Ala Val Pro Pro Gly Pro 65 70 75 80
35	Asn
	(2) INFORMATION FOR SEQ ID NO: 533:
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 70 amino acids (B) TYPE: amino acid
•	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 533:
45	Gly Pro Met Gly Gly Leu Gly Gly Met Glu Ser His His Met Asn Gly 1 5 10 15
50	Ser Leu Gly Ser Gly Asp Met Asp Ser Ile Ser Lys Asn Ser Pro Asn 20 25 30
	Asn Met Ser Leu Ser Asn Gln Pro Gly Thr Pro Arg Asp Asp Gly Glu 35 40 45
55	Met Gly Gly Asn Phe Leu Asn Pro Phe Gln Ser Glu Ser Tyr Ser Pro
60	Ser Met Thr Met Ser Val 65 70

	(2) INFORMATION FOR SEQ ID NO: 534:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 14 amino acids
10	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 534: Thr Cys Glu His Ser Ser Glu Ala Lys Ala Phe His Asp Tyr 1 5 10
15	(2) INFORMATION FOR SEQ ID NO: 535:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 59 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 535:
25	Gln Ala Phe Val Leu Leu Ser Asp Leu Leu Leu Ile Phe Ser Pro Gln 1 5 10 15
	Met Ile Val Gly Gly Arg Asp Phe Leu Arg Pro Leu Val Phe Phe Pro 20 25 30
30	Glu Ala Thr Leu Gln Ser Glu Leu Ala Ser Phe Leu Met Asp His Val 35 40 45
35	Phe Ile Gln Pro Gly Asp Leu Gly Ser Gly Ala 50 55
	(2) INFORMATION FOR SEQ ID NO: 536:
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid
45	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 536: Ala Cys Ser Tyr Leu Leu Cys Asn Pro Glu Phe Thr Phe Phe Ser Arg 1 5 10 15
50	Ala Asp Phe Ala Arg Ser Gln Leu Val Asp Leu Leu Thr Asp Arg Phe
	Gln Gln Glu Leu Glu Leu Leu Gln Val Gly 35 40
55	
	(2) INFORMATION FOR SEQ ID NO: 537:
60	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 35 amino acids

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```
(B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 537:
     Gln Lys Gln Leu Ser Ser Leu Arg Asp Arg Met Val Ala Phe Cys Glu
                                          10
      Leu Cys Gln Ser Cys Leu Ser Asp Val Asp Thr Glu Ile Gln Glu Gln
                                       25
10
      Val Ser Thr
              35
15
      (2) INFORMATION FOR SEQ ID NO: 538:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
20
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 538:
      Gln Val Ile Leu Pro Ala Leu Thr Leu Val Tyr Phe Ser Ile Leu Trp
25
      Thr Leu Thr His Ile Ser Lys Ser Asp Ala Ser
                                       25
                   20
30
       (2) INFORMATION FOR SEQ ID NO: 539:
              (i) SEQUENCE CHARACTERISTICS:
35
                     (A) LENGTH: 31 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 539:
       Ser Thr His Asp Leu Thr Arg Trp Glu Leu Tyr Glu Pro Cys Cys Gln
40
       Leu Leu Gln Lys Ala Val Asp Thr Gly Xaa Val Pro His Gln Val
                                        25
                   20
 45
       (2) INFORMATION FOR SEQ ID NO: 540:
 50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 106 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 540:
 55
       Leu Ala Val Ser Thr Ser Phe Ile Cys Cys Ala Asp Ile Ser Thr Ala
       Leu Pro Leu Gly Ser Ser Arg Pro Ala Pro Ala Pro Arg His Arg Glu
 60
                                      25
                    20
```

	His	Glu	His 35	Gly	His	Gln i	Ala	Arg 40	Pro	Pro	Arg	Leu	Leu 45	Xaa	Thr	Ser
5	Leu	Met 50	Pro	Leu	Ser	Thr	Pro 55	Ala	Ala	Ala	Gln	Leu 60	Leu	Trp	Thr	Gln
10	Leu 65	Thr	Pro	Met	Gly	Gly . 70	Arg	Pro	Gly	Gly	Arg 75	His	Ser	Pro	Pro	Thr 80
	Leu	His	Thr	Gly	Pro 85	Arg	Ala	Leu	Pro	Pro 90	Gly	Pro	Pro	His	Pro 95	Ser
15	Leu	His	Val	Ala 100	Ala	Leu	Ser	Leu	Leu 105	Arg						
20	(2)	INF		(ENCE A) L B) T	CHAI ENGTI YPE :	RACT H: 2 ami	ERIS 07 a	TICS mino		.ds					
25				SEQ		E DE	SCRI	PTIO	N:S							
	Glu 1		ı Val	Leu	Ala 5	Leu	Leu	Trp	Pro	Arg 10		Glu	Leu	Ile	Leu 15	
30	Met	. Ası	n Val	l Gln 20		Val	Arg	Ser	Thr 25		Pro	Gln	Arg	Leu 30		Gly
35	Lev	ı Ası	7hi 3!	r Arg	Pro	His	Tyr	11e 40		Arg	Arg	Tyr	Ala 45		Phe	Ser
	Ser	Ala 5		u Val	. Ser	Ile	Asn 55		Thr	Ile	Pro	Asn 60		Arg	Thr	Met
40	Glr 69		u Le	u Gly	/ Gln	Leu 70		val	. Glu	val	1 G1v 75		Phe	e Val	. Leu	Arg 80
	Va.	l Al	a Al	a Glu	Phe 85		Ser	Arg	, Lys	61\ 90		. Leu	(Val	Phe	Leu 95	Ile
45	Ası	n As	n Ty	r Ası 100		Met	Leu	ı Gly	/ Val		ı Met	. Glu	Arg	Ala 110		Asp
50	As	p Se	r. Ly 11		ı Val	. Glu	Ser	2 Phe		ı Glı	n Let	ı Lev	125		a Arg	Thr
50	Gli	n Gl 13		e Il	e Glu	ı Glu	139		ı Se	r Pr	o Pro	Phe 140		y Gly	Let	ı Val
55	Al 14		e Va	l Ly	s Glu	150		u Ala	a Le	u Il	e Gl: 15:		g Gl	y Gli	n Ala	a Glu 160
	Ar	g Le	n Ar	g Gl	y Gl		ı Al	a Ar	g Va	1 Th 17		n Le	ıIl	e Ar	g Gly 17	y Phe 5
60	G1	v Se	r Se	er Tr	o Lv:	s Sei	Se	r Va	1 G1	u Se	r Le	u Se	r Gl	n As	p Va	l Met

		180		1	85		190	
5	Arg Ser Phe 195		Phe Ar	g Asn G 200	ly Thr S	er Ile Ile 209	e Gln Gly	
10	(2) INFORMA	SEQUENCE (A) (B)	E CHARAC	TERISTI 110 ami	CS: ino acid id	s		
15	(xi) Ala Leu Leu 1	SEQUEN	CE DESCR	RIPTION	SEQ ID	NO: 542: Phe Leu Le	u Gly Asn 15	Glu
20	Arg Ala Th	c Ala Ly 20	s Glu Il	le Arg 1	Asp Glu 1 25	Tyr Val Gl	u Thr Leu 30	Ser
	Lys Ile Tyr		r Tyr Ty	/r Arg 5	Ser Tyr	Leu Gly Ar 4	g Leu Met 5	Lys
25	Val Gln Ty	r Glu Gl		la Glu 1 55	Lys Asp	Asp Leu Me 60	t Gly Val	Glu
30	Asp Thr Al	a Lys Ly	s Gly Pi 70	he Xaa	Ser Lys	Pro Ser Le 75	u Arg Ser	Arg 80
	Asn Thr Il		r Leu G	ly Thr	Arg Gly 90	Ser Val II	le Ser Pro 95	
35	Glu Leu Gl	u Ala Pr 100	o Ile L		Pro His 105	Thr Ala G	in Arg 110	٠
40	(2) INFORM	SEQUEN	CE CHARA	ACTERIST	MICS: ino acid	s		
45	(x	(D)	TOPOLO	GY: lin	ear	D NO: 543:		
	Glu Gln A	rg Tyr P	ro Phe G 5	Slu Ala	Leu Phe 10	Arg Ser G	ln His Tyr 15	
50	Leu Leu A	sp Asn S 20	er Cys 1	Arg Glu	Tyr Leu 25	Phe Ile C	ys Glu Phe 30	≥ Phe
55	:	35		40		Phe His A Asp Ser 1	45	
60		sp Ala I	le Ala '		Leu Cys	Ile His 1	(le Val Le	u Arg

```
Phe Arg Asn Ile Ala Ala Lys Arg Asp Val Pro Ala Leu Asp Arg Tyr
                                         90
                      85
5
     Trp
     (2) INFORMATION FOR SEQ ID NO: 544:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
15
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 544:
     Gly Gly Leu Asp Thr Arg Pro His Tyr Ile Thr Arg Arg Tyr Ala Glu
                                         10
20
      Phe Ser Ser Ala Leu Val Ser Ile Asn Gln
                 20
25
      (2) INFORMATION FOR SEQ ID NO: 545:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 20 amino acids
                    (B) TYPE: amino acid
30
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 545:
      Ser Arg Lys Glu Gln Leu Val Phe Leu Ile Asn Asn Tyr Asp Met Met
                                          10
35
                5
      Leu Gly Val Leu
40
       (2) INFORMATION FOR SEQ ID NO: 546:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 411 amino acids
 45
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 546:
       Ala Leu Leu Lys Tyr Arg Phe Phe Tyr Gln Phe Leu Leu Gly Asn Glu
 50
                                          10
                       5
       Arg Ala Thr Ala Lys Glu Ile Arg Asp Glu Tyr Val Glu Thr Leu Ser
 55
       Lys Ile Tyr Leu Ser Tyr Tyr Arg Ser Tyr Leu Gly Arg Leu Met Lys
                                    40
       Val Gln Tyr Glu Glu Val Ala Glu Lys Asp Asp Leu Met Gly Val Glu
 60
                                                   60
                                55
```

	Asp Thr Ala Lys Lys Gly Phe Xaa Ser Lys Pro Ser Leu Arg Ser Arg 65 70 75 81
5	Asn Thr Ile Phe Thr Leu Gly Thr Arg Gly Ser Val Ile Ser Pro Thr 85 90 95
	Glu Leu Glu Ala Pro Ile Leu Val Pro His Thr Ala Glm Arg Zaa Glu 100 105 110
10	Gln Arg Tyr Pro Phe Glu Ala Leu Phe Arg Ser Gln His Tyr Xma Leu 115 120 125
15	Leu Asp Asn Ser Cys Arg Glu Tyr Leu Phe Ile Cys Glu Phe Phe Val 130 135 140
	Val Ser Gly Pro Xaa Ala His Asp Leu Phe His Ala Val Met Gly 25 145 150 155 150
20	Thr Leu Ser Met Thr Leu Lys His Leu Asp Ser Tyr Leu Ala Asp C/s 165 170 175
	Tyr Asp Ala Ile Ala Val Phe Leu Cys Ile His Ile Val Leu Arg Phe 180 185 130
25	Arg Asn Ile Ala Ala Lys Arg Asp Val Pro Ala Leu Asp Arg Typ Tap 195 200 205
30	Glu Gln Val Leu Ala Leu Leu Trp Pro Arg Phe Glu Leu Ile Leu Glu 210 215 220
	Met Asn Val Gln Ser Val Arg Ser Thr Asp Pro Gln Arg Leu Gly Gly 225 230 235 245
35	Leu Asp Thr Arg Pro His Tyr Ile Thr Arg Arg Tyr Ala Glu Fhe Ser 245 250 255
40	Ser Ala Leu Val Ser Ile Asn Gln Thr Ile Pro Asn Glu Arg Thr Met 260 265 170
40	Gln Leu Leu Gly Gln Leu Gln Val Glu Val Glu Asn Phe Val Leu Arg 275 280 285
45	Val Ala Ala Glu Phe Ser Ser Arg Lys Glu Glm Leu Val Phe Leu Ile 290 295 330
	Asn Asn Tyr Asp Met Met Leu Gly Val Leu Met Glu Arg Ala Ala Asp 305 310 315 320
50	Asp Ser Lys Glu Val Glu Ser Phe Gln Gln Leu Leu Asm Ala Arg Thr 325 330 335
	Gln Glu Phe Ile Glu Glu Leu Leu Ser Pro Pro Pre Gly Gly Leu Val 340 345 350
55	Ala Phe Val Lys Glu Ala Glu Ala Leu Ile Glu Arg Gly Gln Ala Glu 355 360 365
60	Arg Leu Arg Gly Glu Glu Ala Arg Val Thr Glm Leu Ile Arg Gly Phe

	385	Ser	TIP I		390	per /	/aı (siu .		395	JCI (3111 6	ωp (100
5	Arg Ser	Phe		Asn I 105	Phe 1	Arg 1	Asn (Thr 9	Ser					
10	(2) INF		ION I												
15			(A	LE TY () TY	NGTH PE: POLC	: 30 amin XGY:	3 am o ac line	ino id ar	acid		547	:			
20	Tyr Glu	ı Gly	Lys (Glu 5	Phe	Asp	Tyr '	Val	Phe 10	Ser	Ile	Asp '	Val .	Asn 15	Glu
	Gly Gly	y Pro	Ser	Tyr	Lys	Leu	Pro	Tyr 25	Asn	Thr	Ser	Asp	Asp 30	Pro	Trp
25	Leu Th	r Ala 35	Tyr	Asn	Phe	Leu	Gln 40	Lys	Asn	Asp	Leu	Asn 45	Pro	Met	Phe
	Leu As		Val	Ala	Lys	Phe 55	Ile	Ile	Asp	Asn	Thr 60	Lys	Gly	Gln	Met
30	Leu Gl 65	y Leu	Gly	Asn	Pro 70	Ser	Phe	Ser	Asp	Pro 75	Phe	Thr	Gly	Gly	Gly 80
35	Arg Ty	r Val	Pro	Gly 85	Ser	Ser	Gly	Ser	Ser 90	Asn	Thr	Leu	Pro	Th:: 95	Ala
33	Asp Pr	o Phe	Thr 100	Gly	Ala	Gly	Arg	Тут 105	Val	Pro	Gly	Ser	Ala 110	Ser	Met
40	Gly Th	r Thr 115		Ala	Gly	Val	Asp 120	Pro	Phe	Thr	Gly	Asn 125	Ser	Ala	Tyr
	Arg Se		Ala	Ser	Lys	Thr 135	Met	Asn	Ile	Tyr	Phe 140	Pro	Lys	Lys	Glu
45	Ala Va 145	il Thr	Phe	Asp	Gln 150		Asn	Pro	Thr	Gln 155	Ile	Leu	Gly	Lys	Leu 160
50	Lys G	lu Leu	ı Asn	Gly 165		Ala	Pro	Glu	Glu 170		Lys	Leu	Thr	Glu 175	Asp
50	Asp Le	eu Ile	Leu 180		Glu	Lys	Ile	Leu 185		Leu	Ile	Cys	Asn 190		Ser
55	Ser G	lu Ly: 19:		Thr	Val	. Gln	Gln 200		Gln	Ile	Leu	Trp 205		Ala	Ile
	Asn C	ys Pro 10	o Glu	Asp	Ile	val 215		Pro	Ala	Leu	220		Leu	Arg	Leu
60							.			Dha	. ~				. G3.,

,	225	230	235	240
5		Ser Ser His Leu : 245	Ile Asn Leu Leu Ası 250	n Pro Lys Gly 255
	Lys Pro Ala Asn 0		Leu Arg Thr Phe Cy 265	s Asn Cys Phe 270
10	Val Gly Gln Ala (275	Gly Gln Lys Leu 1 280	Met Met Ser Gln Ar 28	g Glu Ser Leu 5
	Met Ser His Ala 290	Ile Glu Leu Lys 295	Ser Gly Ser Asn Ly 300	s Asn Ile
15				
	(2) INFORMATION	FOR SEQ ID NO: 5	48:	
20	(2	NCE CHARACTERIST A) LENGTH: 18 am B) TYPE: amino ac D) TOPOLOGY: line	ino acids cid ear	
	(xi) SEQU	JENCE DESCRIPTION	N: SEQ ID NO: 548:	
25	His Ile Ala Leu 1	Ala Thr Leu Ala 5	Leu Asn Tyr Ser Va	al Cys Phe His 15
	Lys Asp			
30				
	(2) INFORMATION	FOR SEQ ID NO:	549:	
35	(ENCE CHARACTERIS A) LENGTH: 49 am B) TYPE: amino a (D) TOPOLOGY: lir	nino acids Acid	
40			N: SEQ ID NO: 549:	
40	His Asn Ile Glu 1	Gly Lys Ala Gln 5	Cys Leu Ser Leu I 10	le Ser Thr Ile 15
45	Leu Glu Val Val 20		Ala Thr Phe Arg I 25	eu Leu Val Ala 30
	Leu Gly Thr Leu 35	Ile Ser Asp Asp 40	o Ser Asn Ala Val G)	Sln Leu Ala Lys 45
50	Ser			
EE			550.	
5 5		N FOR SEQ ID NO:		
	(i) SEQ	UENCE CHARACTERI (A) LENGTH: 30 a	STICS: mino acids	
		(B) TYPE: amino	acid	
60		(D) TOPOLOGY: 1i	MAST	

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 550:															
5	Leu (_			5					10					Glu 15	Pro
10				20					25					30		
	(2)				FOR :											
15		((xi)	(I	A) LE B) TY D) TO JENCE	PE:	amir OGY:	no ac line	id ear			551	L:			
20	Tyr 1	Glu	Gly	Lys	Glu 5	Phe	Asp	Tyr	Val	Phe 10	Ser	Ile	Asp	Val	Asn 15	Glu
	Gly	Gly	Pro	Ser 20	Tyr	Lys	Leu	Pro	Tyr 25	Asn	Thr	Ser	Asp	Asp 30	Pro	Trp
25	Leu	Thr	Ala 35	Tyr	Asn	Phe	Leu	Gln 40	Lys	Asn	Asp	Leu	Asn 45	Pro	Met	Phe
30	Leu	Asp 50	Gln	Val	Ala	Lys	Phe 55	Ile	Ile	Asp	Asn	Thr 60	Lys	Gly	Gln	Met
30	Leu 65	Gly	Leu	Gly	Asn	Pro 70	Ser	Phe	Ser	Asp	Pro 75	Phe	Thr	Gly	Gly	61y 80
35	Arg	Туг	Val	Pro	Gly 85		Ser	Gly	Ser	Ser 90	Asn	Thr	Leu	Pro	Thr 95	Ala
	Asp	Pro	Phe	Thr 100	Gly	Ala	GĴĀ	Arg	Tyr 105		Pro	Gly	Ser	Ala 110		Met
40	Gly	Thr	Thr 115		Ala	Gly	Val	Asp 120		Phe	Thr	Gly	Asn 125		Ala	Tyr
45	Arg	Ser 130		Ala	Ser	Lys	Thr 135		Asn	Ile	Туг	Phe 140		Lys	Lys	Glu
	Ala 145		Thr	Phe	Asp	Gln 150		Asn	Pro	Thr	Gln 155	Ile	Leu	Gly	Lys	160
50	Lys	Glu	Leu	Asn	Gly 165		Ala	Pro	Glu	Glu 170		Lys	Leu	Thr	179	
	Ąsp	Leu	ılle	180	Leu	Glu	Lys	Ile	185		Leu	Ile	Cys	190		r Ser
55	Ser	Glu	195		Thr	' Val	Gln	Glr 200		Glm	Ile	Leu	205		s Ala	a Ile
	Asn	Cys 210		Glu	ı Asp	Ile	• Val		e Pro	Ala	Leu	Asp 220		e Lei	ı Ar	g Lei

•	Ser Ile 225	Lys I	His		Ser 230	Val	Asn	GIU	AST	235	Cys	ASII	Giu	гÀ2	240
5	Gly Ala	Gln 1	Phe	Ser 245	Ser	His	Leu	Ile	Asn 250	Leu	Leu	Asn	Pro	Lys 255	Gly
	Lys Pro		Asn 260	Gln	Leu	Leu	Ala	Leu 265	Arg	Thr	Phe	Cys	Asn 270	Cys	Phe
10	Val Gly	Gln . 275	Ala	Gly	Gln	Lys	Leu 280	Met	Met	Ser	Gln	Arg 285	Glu	Ser	Leu
15	Met Ser 290	His	Ala	Ile	Glu	Leu 295	Lys	Ser	Gly	Ser	Asn 300	Lys	Asn	Ile	His
15	Ile Ala 305	Leu	Ala	Thr	Leu 310	Ala	Leu	Asn	Tyr	Ser 315	Val	Cys	Phe	His	Lys 320
20	Asp His	Asn	Ile	Glu 325		Lys	Ala	Gln	Cys 330		Ser	Leu	Ile	Ser 335	Thr
	Ile Leu	Glu	Val 340	Val	Gln	Asp	Leu	Glu 345		Thr	Phe	Arg	350	Leu	Val
25	Ala Leu	Gly 355	Thr	Leu	Ile	Ser	360		Ser	Asn	Ala	Val 365	. Glm	. Lev	Ala
30	Lys Ser 370		Gly	Val	Asp	Ser 375		ıle	. Lys	. Lys	Tyr 380	Ser	Ser	· Val	. Ser
•	Glu Pro 385	Ala	Lys	Val	. Ser 390		ı Cys	Cys	Arg	395		. Lev	ı Asr	Le.	400
35															
40	(2) INE	FORMA	TIO	ı FOI	R SE(Q ID	NO:	552	:						
40		(i)	SEQ	(A)	LENG	TH:	TERI 139 ino	amin	o ac	ids					
45		(xi)) SE	(D)	TOPO	LOGY	: li	near	•	ID N	0: 5	52:			
	Tyr Pro	o Asr	ı Gl		p G1; 5	y As	p Il	e Le	u Ar 1		p Gl	n Va	l Le	u Hi 1	s Glu 5
50	His Il	e Glr	n Ar		u Se	r Ly	rs Va		1 Th 5	r Al	a As	n Hi		g Al O	a Leu
55	Gln Il	e Pro		u Va	1 ту	r Le		g G1 10	u Al	a Pr	o Tr		o Se IS	r Al	a Gln
J	Ser Gl 5	u Ile 0	e Ar	g Th	r Il	_	er Al	а Ту	r Ly	rs Th		o A1 0	g As	p Ly	ys Val
60	Gln Cy 65	rs Il	e Le	u Ar		t C) 10	∕a Se	er Th	ır II		t As	n Le	eu Le	eu Se	er Leu 80

	Ala Asn Glu Asp Ser Val Pro Gly Ala Asp Asp Phe Val Pro Val Leu 85 90 95
5	Val Phe Val Leu Ile Lys Ala Asn Pro Pro Cys Leu Leu Ser Thr Val 100 105 110
10	Gln Tyr Ile Ser Ser Phe Tyr Ala Ser Cys Leu Ser Gly Glu Glu Ser 115 120 125
10	Tyr Trp Trp Met Gln Phe Thr Ala Ala Val Glu 130 135
15	(2) INFORMATION FOR SEQ ID NO: 553:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 144 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 553:
25	Tyr Pro Asn Gln Asp Gly Asp Ile Leu Arg Asp Gln Val Leu His Glu 1 5 10 15
	His Ile Gln Arg Leu Ser Lys Val Val Thr Ala Asn His Arg Ala Leu 20 25 30
30	Gln Ile Pro Glu Val Tyr Leu Arg Glu Ala Pro Trp Pro Ser Ala Gln 35 40 45
35	Ser Glu Ile Arg Thr Ile Ser Ala Tyr Lys Thr Pro Arg Asp Lys Val 50 55 60
	Gln Cys Ile Leu Arg Met Cys Ser Thr Ile Met Asn Leu Leu Ser Leu 65 70 75 80
40	Ala Asn Glu Asp Ser Val Pro Gly Ala Asp Asp Phe Val Pro Val Leu 85 90 95
	Val Phe Val Leu Ile Lys Ala Asn Pro Pro Cys Leu Leu Ser Thr Val 100 105 110
45	Gln Tyr Ile Ser Ser Phe Tyr Ala Ser Cys Leu Ser Gly Glu Glu Ser 115 120 125
50	Tyr Trp Trp Met Gln Phe Thr Ala Ala Val Glu Phe Ile Lys Thr Ile 130 135 140
55	(2) INFORMATION FOR SEQ ID NO: 554:
	(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 14 amino acids
(B) TYPE: amino acid

```
(D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 554:
     Tyr Pro Asn Gln Asp Gly Asp Ile Leu Arg Asp Gln Val Leu
 5
                        5
       1
      (2) INFORMATION FOR SEQ ID NO: 555:
10
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 11 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 555:
15
      Glu Ala Pro Trp Pro Ser Ala Gln Ser Glu Ile
                        5
20
      (2) INFORMATION FOR SEQ ID NO: 556:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 21 amino acids
25
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 556:
      Ser Gly Glu Glu Ser Tyr Trp Trp Met Gln Phe Thr Ala Ala Val Glu
30
      Phe Ile Lys Thr Ile
                   20
35
       (2) INFORMATION FOR SEQ ID NO: 557:
 40
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 18 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 557:
 45
       Ala Asp Asp Phe Val Pro Val Leu Val Phe Val Leu Ile Lys Ala Asn
                        5
                                            10
       Pro Pro
 50
       (2) INFORMATION FOR SEQ ID NO: 558:
. 55
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 12 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
 60
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 558:
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Tyr Lys Thr Pro Arg Asp Lys Val Gln Cys Ile Leu
                       5
5
      (2) INFORMATION FOR SEQ ID NO: 559:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 15 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 559:
      Gly Ala Asp Asp Phe Val Pro Val Leu Val Phe Val Leu Ile Lys
15
                        5
20
      (2) INFORMATION FOR SEQ ID NO: 560:
              (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 12 amino acids
                    (B) TYPE: amino acid
25
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 560:
      Pro Val Leu Val Phe Val Leu Ile Lys Ala Asn Pro
                                           10
                        5
        1
30
      (2) INFORMATION FOR SEQ ID NO: 561:
35
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 17 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 561:
40
      Ser Ala Arg Ala Ser Thr Gln Pro Pro Ala Gly Gln His Pro Gly Pro
        1
                                            10
      Cys
45
       (2) INFORMATION FOR SEQ ID NO: 562:
 50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 33 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
 55
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 562:
       Met Pro Gly Arg Trp Arg Trp Gln Arg Asp Met His Pro Ala Arg Lys
                                                                15 .
                                            10
       Leu Leu Ser Leu Leu Phe Leu Ile Leu Met Gly Thr Glu Leu Thr Gln
 60
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30 20 25 Asp 5 (2) INFORMATION FOR SEQ ID NO: 563: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 19 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 563: 15 Ser Ala Ala Pro Asp Ser Leu Leu Arg Ser Ser Lys Gly Ser Thr Arg 10 5 1 Gly Ser Leu 20 (2) INFORMATION FOR SEQ ID NO: 564: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 564: 30 Ala Ala Ile Val Ile Trp Arg Gly Lys Ser Glu Ser Arg Ile Ala Lys 10 35 Thr Pro Gly Ile 20 40 (2) INFORMATION FOR SEQ ID NO: 565: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 17 amino acids (B) TYPE: amino acid 45 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 565: Pro Leu Gly Ile Thr Leu Pro Leu Gly Ala Pro Glu Thr Gly Gly 10 5 50 Asp 55 (2) INFORMATION FOR SEQ ID NO: 566: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids

(B) TYPE: amino acid

```
(D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 566:
     Cys Ala Ala Glu Thr Trp Lys Gly Ser Gln Arg Ala Gly Gln Leu Cys
                                          10
 5
     Ala Leu Leu Ala
                  20
10
      (2) INFORMATION FOR SEQ ID NO: 567:
             (i) SEQUENCE CHARACTERISTICS:
15
                    (A) LENGTH: 20 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 567:
      Phe Arg Gly Gly Gly Thr Leu Val Leu Pro Pro Thr His Thr Pro Glu
20
                                          10
      Trp Leu Ile Leu
                   20
25
      (2) INFORMATION FOR SEQ ID NO: 568:
              (i) SEQUENCE CHARACTERISTICS:
30
                     (A) LENGTH: 22 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 568:
35
      Met Arg Ser Ala Arg Pro Ser Leu Gly Cys Leu Pro Ser Trp Ala Phe
       Ser Gln Ala Leu Asn Ile
 40
                    20
       (2) INFORMATION FOR SEQ ID NO: 569:
 45
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 22 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 569:
 50
       Leu Leu Gly Leu Lys Gly Leu Ala Pro Ala Glu Ile Ser Ala Val Cys
       Glu Lys Gly Asn Phe Asn
 55
                    20
 60
       (2) INFORMATION FOR SEQ ID NO: 570:
```

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 570:
     Val Ala His Gly Leu Ala Trp Ser Tyr Tyr Ile Gly Tyr Leu Arg Leu
                                          10
10
     Ile Leu Pro Glu Leu Gln Ala Arg Ile Arg
                  20
15
      (2) INFORMATION FOR SEQ ID NO: 571:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 18 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 571:
      Thr Tyr Asn Gln His Tyr Asn Asn Leu Leu Arg Gly Ala Val Ser Gln
25
                                           10
      Arg Cys
30
      (2) INFORMATION FOR SEQ ID NO: 572:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 43 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 572:
40
      Ile Leu Leu Pro Leu Asp Cys Gly Val Pro Asp Asn Leu Ser Met Ala
                                           10
                        5
        1
      Asp Pro Asn Ile Arg Phe Leu Asp Lys Leu Pro Gln Gln Thr Gly Asp
                                       25
45
      Arg Ala Gly Ile Lys Asp Arg Val Tyr Ser Asn
                                   40
 50
       (2) INFORMATION FOR SEQ ID NO: 573:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 45 amino acids
 55
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 573:
       Ser Ile Tyr Glu Leu Leu Glu Asn Gly Gln Arg Ala Gly Thr Cys Val
 60
                         5
                                           10
```

	Leu Glu Tyr Ala Thr Pro Leu Gln Thr Leu Phe Ala Met Ser Gln Tyr 20 25 30
5	Ser Gln Ala Gly Phe Ser Gly Glu Asp Arg Leu Glu Gln 35 40 45
10	(2) INFORMATION FOR SEQ ID NO: 574:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids
15	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 574:
20	Ala Lys Leu Phe Cys Arg Thr Leu Glu Asp Ile Leu Ala Asp Ala Pro 1 5 10 15
20	Glu Ser Gln Asn Asn Cys Arg Leu Ile Ala Tyr Gln Glu Pro Ala Asp 20 25 30
25	Asp Ser Ser Phe Ser Leu Ser Gln Glu Val Leu Arg His Leu Arg Gln 35 40 45
	Glu Glu Lys Glu Glu Val Thr Val Gly Ser Leu Lys Thr Ser Ala Val 50 55 60
30	Pro Ser Thr Ser Thr Met Ser Gln Glu Pro Glu Leu Leu Ile Ser Gly 65 70 75 80
	Met Glu Lys Pro Leu Pro Leu Arg Thr Asp Phe Ser 85 90
35	
	(2) INFORMATION FOR SEQ ID NO: 575:
40	(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 43 amino acids(B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 575:
45	Leu Leu Gly Leu Lys Gly Leu Ala Pro Ala Glu Ile Ser Ala Val Cy 1 5 10 15
50	Glu Lys Gly Asn Phe Asn Val Ala His Gly Leu Ala Trp Ser Tyr Ty 20 25 30
	Ile Gly Tyr Leu Arg Leu Ile Leu Pro Glu Leu 35 40
55	
	(2) INFORMATION FOR SEQ ID NO: 576:
	(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 22 amino acids

```
(3) T/PE: amino acid
                    (D) TCPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 576:
      Thr Met Lys Leu Lys Leu Arg Arg Asn Ile Val Lys Leu Ser Leu
                       5
                                          10
      Tyr Arg His The Thr Asn
                  20
10
      (2) DEFORMATION FOR SEQ ID NO: 577:
15
             (1) SEQUENCE CHARACTERISTICS:
                    (A) LEXGTH: 22 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 577:
20
     Thr Leu Ile Leu Ala Val Ala Ala Ser Ile Val Phe Ile Ile Trp Thr
                       5
                                          10
     Thr Met Lys Phe Arg Ile
25
                  7:
      (2) DEFORMATION FOR SEQ ID NO: 578:
30
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 28 amino acids
                     '3) TYPE: amino acid
                     D, TOPOLOGY: linear
35
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 578:
     Val Thr Cys Gim Ser Asp Trp Arg Glu Leu Trp Val Asp Asp Ala Ile
40
     Trp Arg Leu Leu Phe Ser Met Ile Leu Phe Val Ile
                  20
45
      (2) DEFORMATION FOR SEQ ID NO: 579:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
                    (3) TYPE: amino acid
50
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 579:
     Met Val Leu Trp Arg Pro Ser Ala Asn Asn Gln Arg Phe Ala Phe Ser
                       5
                                          10
55
     Pro Leu Ser Glu Glu Glu Glu Glu Asp Glu Gln
```

```
(2) INFORMATION FOR SEQ ID NO: 580:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 27 amino acids
 5
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 580:
      Met Val Leu Trp Arg Pro Ser Ala Asn Asn Gln Arg Phe Ala Phe Ser
10
      Pro Leu Ser Glu Glu Glu Glu Glu Asp Glu Gln
                   20
15
      (2) INFORMATION FOR SEQ ID NO: 581:
              (i) SEQUENCE CHARACTERISTICS:
20
                     (A) LENGTH: 35 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 581:
25
      Lys Glu Pro Met Leu Lys Glu Ser Phe Glu Gly Met Lys Met Arg Ser
        1
                        5
      Thr Lys Gln Glu Pro Asn Gly Asn Ser Lys Val Asn Lys Ala Gln Glu
30
      Asp Asp Leu
               35
35
      (2) INFORMATION FOR SEQ ID NO: 582:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 37 amino acids
40
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 582:
      Lys Trp Val Glu Glu Asn Val Pro Ser Ser Val Thr Asp Val Ala Leu
45
                                           10
      Pro Ala Leu Leu Asp Ser Asp Glu Glu Arg Met Ile Thr His Phe Glu
                                      25
50
      Arg Ser Lys Met Glu
               35
55
      (2) INFORMATION FOR SEQ ID NO: 583:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 20 amino acids
                     (B) TYPE: amino acid
60
                    (D) TOPOLOGY: linear
```

```
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 583:
     Asp Pro Arg Val Arg Leu Asn Ser Leu Thr Cys Lys His Ile Phe Ile
1 5 11 15
 5
     Ser Leu Thr Gin
10
      (2) INFORMATION FOR SEQ ID NO: E34:
             (i) SEQUENCE CHAPACTERISTICS:
                    (A) LENGTH: 13 amuno aciis
                     (B) TYFE: amino acid
15
                    (D) TOPOLOTY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 584:
     Tyr Glu Pro Met Asp Phe Yaa Met Ala Le: Ile Tyr Asp
20
      (2) INFORMATION FOR SEQ ID NO: 535:
25
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 18 amino acids
                     (B) TYFE: amimo acid
                     (D) TOPOLOGY: Limear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 515:
      Ile Arg His Glu Leu Thr Val Leu Arg Asp Thr Arg Fro Ala Dys Ala
35
40
      (2) INFORMATION FOR SEQ ID NO: ESE:
              (i) SEQUENCE THAFACTERISTICS:
                     (A) LENGTH: 10 amins estis
                     (B) TiFE: amino asid
45
                     (D) TCPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 506:
      Met Asp Phe Xaa Ket Ala Leu Ile Tyr Asp
 50
       (2) INFORMATION FOR SEQ ID NO: 537:
55
              (i) SEQUENCE CHAFACTERISTICS:
                     (A) LENGTH: 14 amino acids
                     (B) T:PE: amino acid
                     (D) TCPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 557:
 60
```

1 5 10 15
Ser Ile Pro Gly Gly Tyr Asn Ala 20
(2) INFORMATION FOR SEQ ID NO: 588: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 25 amino acids
(A) LENGTH: 25 Antho acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 588:
Leu Arg Arg Met Tyr Thr Asp Ile Gln Glu Pro Met Leu Ser Ala Ala 1 5 10 15
Gln Glu Gln Phe Gly Gly Asn Pro Phe 20 25
(2) INFORMATION FOR SEQ ID NO: 589:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 32 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 589:
Ala Ser Leu Val Ser Asn Thr Ser Ser Gly Glu Gly Ser Gln Pro Ser 1 5 10 15
Arg Thr Glu Asn Arg Asp Pro Leu Pro Asn Pro Trp Ala Pro Gln Thr 20 25 30
(2) INFORMATION FOR SEQ ID NO: 590: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 71 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 590:
Ser Gln Ser Ser Ser Ala Ser Ser Gly Thr Ala Ser Thr Val Gly Gly 1 5 10 15
Thr Thr Gly Ser Thr Ala Ser Gly Thr Ser Gly Gln Ser Thr Thr Ala 20 25 30
Pro Asn Leu Val Pro Gly Val Gly Ala Ser Met Phe Asn Thr Pro Gly 35 40 45

	Met Gln Ser Leu Leu Gln Gln Ile Thr Glu Asn Pro Gln Leu Met Gln 50 55 60
5	Asn Met Leu Ser Ala Pro Tyr 65 70
10	(2) INFORMATION FOR SEQ ID NO: 591: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 45 amino acids
15	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 591:
	Met Arg Ser Met Met Gln Ser Leu Ser Gln Asn Pro Asp Leu Ala Ala 1 5 10 15
20	Gln Met Met Leu Asn Asn Pro Leu Phe Ala Gly Asn Pro Gln Leu Gln 20 25 30
25	Glu Gln Met Arg Gln Gln Leu Pro Thr Phe Leu Gln Gln 35 40 45
30	(2) INFORMATION FOR SEQ ID NO: 592: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 73 amino acids (B) TYPE: amino acid
35	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 592: Met Gln Asn Pro Asp Thr Leu Ser Ala Met Ser Asn Pro Arg Ala Met 1 5 10 15
40	Gln Ala Leu Leu Gln Ile Gln Gln Gly Leu Gln Thr Leu Ala Thr Glu 20 25 30
	Ala Pro Gly Leu Ile Pro Gly Phe Thr Pro Gly Leu Gly Ala Leu Gly 35 40 45
45	Ser Thr Gly Gly Ser Ser Gly Thr Asn Gly Ser Asn Ala Thr Pro Ser 50 55 60
50	Glu Asn Thr Ser Pro Thr Ala Gly Thr 65 70
	(2) INFORMATION FOR SEQ ID NO: 593:
55	(A) LENGTH: 72 amino acids (B) TYPE: amino acid
60	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 593:

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The Glu Peo Gly His Gln Gln Phe Ile Gln Gln Met Leu Gln Ala Leu 10 Ala Sly Val Asn Pro Gln Leu Gln Asn Pro Glu Val Arg Phe Gln Gln 5 Glm Leu Glu Gln Leu Ser Ala Met Gly Phe Leu Asn Arg Glu Ala Asn 40 Let 31m Ala Leu Ile Ala Thr Gly Gly Asp Ile Asn Ala Ala Ile Glu 10 50 Arg Leu Leu Gly Ser Gln Pro Ser 70 15 (2) DESCRIPTION FOR SEQ ID NO: 594: (i) SEQUENCE CHARACTERISTICS: 20 (A) LENGTH: 45 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 594: 25 Arg Asn Pro Ala Met Met Gln Glu Met Met Arg Asn Gln Asp Arg Ala Let Ser Ash Leu Glu Ser Ile Pro Gly Gly Tyr Ash Ala Leu Arg Arg 25 30 Met Tyr Thr Asp Ile Gln Glu Pro Met Leu Ser Ala Ala 40 35 35 (2) DECEMATION FOR SEQ ID NO: 595: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 13 amino acids 40 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 595: Gly Asn Pro Phe Ala Ser Leu Val Ser Asn Thr Ser Ser 45 5 (2) INFORMATION FOR SEQ ID NO: 596: 50 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 11 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 55 (Zi) SEQUENCE DESCRIPTION: SEQ ID NO: 596: Glu Asn Arg Asp Pro Leu Pro Asn Pro Trp Ala 5 60

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(2) INFORMATION FOR SEQ ID NO: 597:
             (i) SEQUENCE CHARACTERISTICS:
5
                    (A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 597:
10
     Gly Lys Ile Leu Lys Asp Gln Asp Thr Leu Ser Gln His Gly Ile His
                                           10
       1
      Asp
15
      (2) INFORMATION FOR SEQ ID NO: 598:
20
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 14 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 598:
25
      Gly Leu Thr Val His Leu Val Ile Lys Thr Gln Asn Arg Pro
                         5
 30
       (2) INFORMATION FOR SEQ ID NO: 599:
              (i) SEQUENCE CHARACTERISTICS:
 35
                      (A) LENGTH: 18 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY; linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 599:
       Ser Glu Leu Gln Ser Gln Met Gln Arg Gln Leu Leu Ser Asn Pro Glu
 40
                                            10
                         5
         1
       Met Met
 45
       (2) INFORMATION FOR SEQ ID NO: 600:
               (i) SEQUENCE CHARACTERISTICS:
 50
                      (A) LENGTH: 14 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 600:
 55
       Pro Glu Ile Ser His Met Leu Asn Asn Pro Asp Ile Met Arg
                                             10
                          5
         1
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(2) INFORMATION FOR SEQ ID NO: 501:
            (i) SEQUENCE CHAPACTERISTICS:
                   (A) LENGTH: 18 amino acids
                    (3) TYPE: amino acid
5
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 601:
     Arg Gln Leu Ile Met Ala Asn Pro Gln Met Gln Gln Leu Ile Gln Arg
                                     10
10
                    5
     Asn Pro
15
      (2) INFORMATION FOR SEQ ID NO: 502:
             (i) SEQUENCE CHAPACTERISTICS:
                    (A) LENGTH: 27 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 502:
      Asn Leu Cys His Val Asp Cys Gln Asp Leu Leu Asn Pro Asn Leu Leu
25
                                  15
      Ala Gly Ile His Cys Ala Lys Arg Ile Val Ser
                                       25
                   20
30
      (2) INFORMATION FOR SEQ ID NO: 503:
35
              (i) SECUTICE CHARACTERISTICS:
                     (A) LENGTH: 23 amino acids
                     (3) TYPE: amino acid
                    (D) TOPOLOGY: limear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 503:
40
      Leu Asp Gly Phe Glu Gly Tyr Ser Leu Ser Asp Trp Leu Cys Leu Ala
                                          10
       Phe Val Glu Ser Lys Phe Asn
 45
                   20
       (2) INFORMATION FOR SEQ ID NO: 604:
 50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LEXGTH: 22 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
 55
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504:
       Asn Glu Asn Ala Asp Gly Ser Phe Asp Tyr Gly Leu Phe Gln Ile Asn
                                           10
 60
       Ser His Tyr Trp Cys Asn
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(2) INFORMATION FOR SEQ ID NO: 605:
            (i) SEQUENCE CHARACTERISTICS:
                   (A) LENGTH: 27 amino acids
                    (B) TYPE: amino acid
                   (D) TOPOLOGY: linear
10
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 605:
     Asn Leu Cys His Val Asp Cys Gln Asp Leu Leu Asn Pro Asn Leu Leu
15
      Ala Gly Ile His Cys Ala Lys Arg Ile Val Ser
                                     25
                  20
20
      (2) INFORMATION FOR SEQ ID NO: 606:
         (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 13 amino acids
                    (B) TYPE: amino acid
25
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 606:
      Ile Arg Glu Val Asn Glu Val Ile Gln Asn Pro Ala Thr
30
                        5
      (2) INFORMATION FOR SEQ ID NO: 607:
 35
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 30 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 607:
 40
       Ile Thr Arg Ile Leu Leu Ser His Phe Asn Trp Asp Lys Glu Lys Leu
                                10
       Met Glu Arg Tyr Phe Asp Gly Asn Leu Glu Lys Leu Phe Ala
 45
                                     25
                    20
       (2) INFORMATION FOR SEQ ID NO: 608:
 50
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 23 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
  55
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 608:
       Asn Thr Arg Ser Ser Ala Gln Asp Met Pro Cys Gln Ile Cys Tyr Leu
                                   10
                 5
  60
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Asn Tyr Pro Asn Ser Tyr Phe
             20
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5 (2) INFORMATION FOR SEQ ID NO: 609: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 60 amino acids (B) TYPE: amino acid 10 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 609: Cys Asp Ile Leu Val Asp Asp Asn Thr Val Met Arg Leu Ile Thr Asp 15 10 Ser Lys Val Lys Leu Lys Tyr Gln His Leu Ile Thr Asn Ser Phe Val Glu Cys Asn Arg Leu Leu Lys Trp Cys Pro Ala Pro Asp Cys His His 20 40 Val Val Lys Val Gln Tyr Pro Asp Ala Lys Pro Val 55 25 (2) INFORMATION FOR SEQ ID NO: 610: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 52 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 610: 35 Cys Asp Ile Leu Val Asp Asp Asn Thr Val Met Arg Leu Ile Thr Asp - · 10 5 Ser Lys Val Lys Leu Lys Tyr Gln His Leu Ile Thr Asn Ser Phe Val 40 Glu Cys Asn Arg Leu Leu Lys Trp Cys Pro Ala Pro Asp Cys His His 40 45 Val Val Lys Val 50 50 (2) INFORMATION FOR SEQ ID NO: 611: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 60 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 611:

60

Gly Cys Asn His Met Val Cys Arg Asn Gln Asn Cys Lys Ala Glu Phé

Cys Trp Val Cys Leu Gly Pro Trp Glu Pro His Gly Ser Ala Trp Tyr 25 Asn Cys Asn Arg Tyr Asn Glu Asp Asp Ala Lys Ala Ala Arg Asp Ala 5 Gln Glu Arg Ser Arg Ala Ala Leu Gln Arg Tyr Leu 55 10 (2) INFORMATION FOR SEQ ID NO: 612: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 60 amino acids 15 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 612: Phe Tyr Cys Asn Arg Tyr Met Asn His Met Gln Ser Leu Arg Phe Glu 20 His Lys Leu Tyr Ala Gln Val Lys Gln Lys Met Glu Glu Met Gln Gln 25 25 His Asn Met Ser Trp Ile Glu Val Gln Phe Leu Lys Lys Ala Val Asp Val Leu Cys Gln Cys Arg Ala Thr Leu Met Tyr Thr 55 30 (2) INFORMATION FOR SEQ ID NO: 613: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 60 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 613: 40 Tyr Val Phe Ala Phe Tyr Leu Lys Lys Asn Asn Gln Ser Ile Ile Phe 10 Glu Asn Asn Gln Ala Asp Leu Glu Asn Ala Thr Glu Val Leu Ser Gly 45 Tyr Leu Glu Arg Asp Ile Ser Gln Asp Ser Leu Gln Asp Ile Lys Gln 40 50 Lys Val Gln Asp Lys Tyr Arg Tyr Cys Glu Ser Arg 55 55 (2) INFORMATION FOR SEQ ID NO: 614: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 37 amino acids (B) TYPE: amino acid 60

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(D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 614:
     Thr Gly Leu Glu Cys Gly His Lys Phe Cys Met Gln Cys Trp Ser Glu
5
     Tyr Leu Thr Thr Lys Ile Met Glu Glu Gly Met Gly Gln Thr Ile Ser
     Cys Pro Ala His Gly
10
              35
      (2) INFORMATION FOR SEQ ID NO: 615:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 21 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
20
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 615:
      Met Trp Gly Tyr Leu Phe Val Asp Ala Ala Trp Asn Phe Leu Gly Cys
                                           10
                        5
25
      Leu Ile Cys Gly Trp
                   20
30
       (2) INFORMATION FOR SEQ ID NO: 616:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 46 amino acids
                     (B) TYPE: amino acid
 35
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 616:
       Met His Phe Ile Ser Ser Gly Asn Val Ser Ala Ile Arg Ser Ser Ile
 40
       Leu Leu Arg Xaa Ser Leu Ser Tyr Leu Gly Asn Cys Leu Arg Val
       Ser Ala Ile Phe Val Tyr Phe Leu Leu Phe Leu Leu Leu Ser
 45
                                     40
  50
        (2) INFORMATION FOR SEQ ID NO: 617:
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 80 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
  55
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 617:
        Met Asp Gln Ala Leu Arg Gly Ser Pro Ser Glu Gly Phe Ser Thr Asp
                          5
  60
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	Pro Ser Pro Pro Gln Val Gly Arg Gln Ile Pro Ser Phe Pro Pro Trp 20 25 30
5	Arg Arg Leu Val Leu Pro Lys Ala Ser Gly Cys Phe Leu Glu Arg Glu 35 40 45
	Trp Trp Leu Cys Val Phe Lys Leu Arg Thr Arg Pro Gly Ala Glu Ala 50 55 60
10	His Ala Tyr Asn Ser Ser Ile Leu Gly Gly Arg Gly Lys Gly Ile Thr 65 70 75 80
15	•
	(2) INFORMATION FOR SEQ ID NO: 618:
20	(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 131 amino acids(B) TYPE: amino acid
25	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 618:
	Met Leu Pro Ala Leu Ala Ser Cys Cys His Phe Ser Pro Pro Glu Gln 1 5 10 15
30	Ala Ala Arg Leu Lys Lys Leu Gln Glu Gln Glu Lys Gln Gln Lys Val 20 25 30
	Glu Phe Arg Lys Arg Met Glu Lys Glu Val Ser Asp Phe Ile Gln Asp 35 40 45
35	Ser Gly Gln Ile Lys Lys Lys Phe Gln Pro Met Asn Lys Ile Glu Arg 50 55 60
40	Ser Ile Leu His Asp Val Val Glu Val Ala Gly Leu Thr Ser Phe Ser 65 70 75 80
	Phe Gly Glu Asp Asp Asp Cys Arg Tyr Val Met Ile Phe Lys Lys Glv 85 90 95
45	Phe Ala Pro Ser Asp Glu Glu Leu Asp Ser Tyr Arg Arg Gly Glu Glu 100 105 110
	Trp Asp Pro Gln Lys Ala Glu Glu Lys Arg Asn Xaa Lys Glu Leu Al 115 120 125
50	Gln Arg Gln 130
55	(2) INFORMATION FOR SEQ ID NO: 619:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 76 amino acids
60	(B) TYPE: amino acid(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 619:
5	Glu Glu Glu Ala Ala Gln Gln Gly Pro Val Val Val Ser Pro Ala Ser 1 5 10 15
,	Asp Tyr Lys Asp Lys Tyr Ser His Leu Ile Gly Lys Gly Ala Ala Lys 20 25 30
10	Asp Ala Ala His Met Leu Gln Ala Asn Lys Thr Tyr Gly Cys Xaa Pro 35 40 45
	Val Ala Asn Lys Arg Asp Thr Arg Ser Ile Glu Glu Ala Met Asn Glu 50 55 60
15	Ile Arg Ala Lys Lys Arg Leu Arg Gln Ser Gly Glu 65 70 75
20	(2) INFORMATION FOR SEQ ID NO: 620:
25	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 40 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 620:
30	Pro Pro Arg Arg Pro Ala Gln Leu Pro Leu Thr Pro Gly Ala Gly Gln 1 5 10 15
50	Gly Ala Gly Arg Asp Lys Ala Ala Ala Ile Arg Ala His Pro Gly Ala 20 25 30
35	Pro Pro Leu Asn His Leu Leu Pro 35 40
40	(2) INFORMATION FOR SEQ ID NO: 621: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 28 amino acids (B) TYPE: amino acid
45	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 621:
	Ala Val Pro Gln Ala Gly Gly Lys Gln Val Phe Asp Leu Ser Pro Leu 1 5 10 15
50	Glu Leu Gly Tyr Val Arg Gly Met Cys Val Cys Val 20 25
55	(2) INFORMATION FOR SEQ ID NO: 622:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 207 amino acids
60	(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 622	(xi)	SEQUENCE	DESCRIPTION:	SEQ	ID NO:	622:
---	------	----------	--------------	-----	--------	------

Met Leu Pro Ala Leu Ala Ser Cys Cys His Phe Ser Pro Pro Glu Gln 10

5

Ala Ala Arg Leu Lys Lys Leu Gln Glu Gln Glu Lys Gln Gln Lys Val

10

Glu Phe Arg Lys Arg Met Glu Lys Glu Val Ser Asp Phe Ile Gln Asp

Ser Gly Gln Ile Lys Lys Phe Gln Pro Met Asn Lys Ile Glu Arg 55

15

Ser Ile Leu His Asp Val Val Glu Val Ala Gly Leu Thr Ser Phe Ser

Phe Gly Glu Asp Asp Cys Arg Tyr Val Met Ile Phe Lys Lys Glu 90

20

Phe Ala Pro Ser Asp Glu Glu Leu Asp Ser Tyr Arg Arg Gly Glu Glu

25

Trp Asp Pro Gln Lys Ala Glu Glu Lys Arg Asn Xaa Lys Glu Leu Ala 120

Gln Arg Gln Glu Glu Glu Ala Ala Gln Gln Gly Pro Val Val Val Ser 135

30

Pro Ala Ser Asp Tyr Lys Asp Lys Tyr Ser His Leu Ile Gly Lys Gly 155

Ala Ala Lys Asp Ala Ala His Met Leu Gln Ala Asn Lys Thr Tyr Gly 170

35

Cys Xaa Pro Val Ala Asn Lys Arg Asp Thr Arg Ser Ile Glu Glu Ala 185 180

Met Asn Glu Ile Arg Ala Lys Lys Arg Leu Arg Gln Ser Gly Glu 200 40

(2) INFORMATION FOR SEQ ID NO: 623:

45

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 34 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 623:

Leu Leu Cys Pro Val Leu Asn Ser Gly Xaa Ser Trp Asn Phe Pro His 10 5

Pro Ser Gln Pro Glu Tyr Ser Phe His Gly Phe His Ser Thr Arg Leu 55

Trp Ile

	(2) INFORMATION FOR SEQ ID NO: 624:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 28 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
0	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 624: Pro Ser Thr Pro Trp Phe Leu Phe Leu Leu Gly Leu Thr Cys Pro Phe
	1 5 10 15
15	Ser Thr Ser His Pro Arg Trp Asp Ser Ile Pro Pro 20 25
20	(2) INFORMATION FOR SEQ ID NO: 625: (i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 227 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 625:
	Glu Leu Ser Ile Ser Ile Ser Asn Val Ala Leu Ala Asp Glu Gly Glu 1 5 10 15
30	Tyr Thr Cys Ser Ile Phe Thr Met Pro Val Arg Thr Ala Lys Ser Leu 20 25 30
35	Val Thr Val Leu Gly Ile Pro Gln Lys Pro Ile Ile Thr Gly Tyr Lys 35 40 45
22	Ser Ser Leu Arg Glu Lys Asp Thr Ala Thr Leu Asn Cys Gln Ser Ser 50 55 60
40	Gly Ser Lys Pro Ala Ala Arg Leu Thr Trp Arg Lys Gly Asp Gln Glu 65 70 75 80
	Leu His Gly Glu Pro Thr Arg Ile Gln Glu Asp Pro Asn Gly Lys Thr 85 90 95
45	Phe Thr Val Ser Ser Ser Val Thr Phe Gln Val Thr Arg Glu Asp Asp 100 105 110
50	Gly Ala Ser Ile Val Cys Ser Val Asn His Glu Ser Leu Lys Gly Ala 115 120 125
50	Asp Arg Ser Thr Ser Gln Arg Ile Glu Val Leu Tyr Thr Pro Thr Ala 130 135 140
55	Met Ile Arg Pro Asp Pro Pro His Pro Arg Glu Gly Gln Lys Leu Leu 145 150 155 160
	Leu His Cys Glu Gly Arg Gly Asn Pro Val Pro Gln Gln Tyr Leu Trp 165 170 175 .
60	Glu Lys Glu Gly Ser Val Pro Pro Leu Lys Met Thr Gln Glu Ser Ala

	180	185	190
	Leu Ile Phe Pro Phe Leu Asr 195	n Lys Ser Asp Ser Gly 200	Thr Tyr Gly Cys 205
5	Thr Ala Thr Ser Asn Met Gly 210 219		Tyr Thr Leu Asn
10	Val Asn Asp 225		
15		TERISTICS: 64 amino acids	
20		<pre>(: linear RIPTION: SEQ ID NO: 6:</pre>	
	Glu Leu Ser Ile Ser Ile Se	er Asn Val Ala Leu Ala 10	a Asp Glu Gly Glu 15
25	Tyr Thr Cys Ser Ile Phe Tr 20	or Met Pro Val Arg Th 25	r Ala Lys Ser Leu 30
30	Val Thr Val Leu Gly Ile Pr 35	co Gln Lys Pro Ile Il 40	e Thr Gly Tyr Lys 45
50	Ser Ser Leu Arg Glu Lys A	sp Thr Ala Thr Leu As 55	n Cys Gln Ser Ser
35			
40	(2) INFORMATION FOR SEQ I (i) SEQUENCE CHARP	CTERISTICS:	
	(B) TYPE: a		
45	(xi) SEQUENCE DESC	CRIPTION: SEQ ID NO:	627 :
	Cys Gln Ser Ser Gly Ser I 1 5	ys Pro Ala Ala Arg L 10	eu Thr Trp Arg Lys 15
50	Gly Asp Gln Glu Leu His (Gly Glu Pro Thr Arg I 25	le Gln Glu Asp Pro 30
e c	Asn Gly Lys Thr Phe Thr 1 35	Val Ser Ser Ser Val T 40	hr Phe Gln Val Thi 45
55	Arg Glu Asp Asp Gly Ala : 50	Ser Ile Val Cys Ser V 55	al Asn His Glu Se 60
60	Leu 65		

e	(2) INFORMATION FOR SEQ ID NO: 628:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 58 amino acids
	(B) TYPE: amino acid
10	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 628:
10	
	His Glu Ser Leu Lys Gly Ala Asp Arg Ser Thr Ser Gln Arg Ile Glu 1 5 10 15
15	Val Leu Tyr Thr Pro Thr Ala Met Ile Arg Pro Asp Pro Pro His Pro 20 25 30
20	Arg Glu Gly Gln Lys Leu Leu His Cys Glu Gly Arg Gly Asn Pro 35 40 45
20	Val Pro Gln Gln Tyr Leu Trp Glu Lys Glu 50 55
25	
23	(2) INFORMATION FOR SEQ ID NO: 629:
	(i) SEQUENCE CHARACTERISTICS:
20	(A) LENGTH: 52 amino acids
30	(B) TYPE: amino acid (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 629:
35	Trp Glu Lys Glu Gly Ser Val Pro Pro Leu Lys Met Thr Gln Glu Ser 1 5 10 15
	Ala Leu Ile Phe Pro Phe Leu Asn Lys Ser Asp Ser Gly Thr Tyr Gly
	20 25 30
40	Cys Thr Ala Thr Ser Asn Met Gly Ser Tyr Lys Ala Tyr Tyr Thr Leu 35 40 45
	Asn Val Asn Asp
45	50
73	
	(2) INFORMATION FOR SEQ ID NO: 630:
50	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 123 amino acids(B) TYPE: amino acid
-	(D) TOPOLOGY: linear
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 630:
J.	Val Pro Glu Leu Pro Asp Arg Val His Gln Leu His Gln Ala Val Gln 1 5 10 15
60	Gly Cys Ala Leu Gly Arg Pro Gly Phe Pro Gly Gly Pro Thr His Ser 20 25 30

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Gly His His Lys Ser His Pro Gly Pro Ala Gly Gly Asp Tyr Asn Arg 40 Cys Asp Arg Pro Gly Gln Val His Leu His Asn Pro Arg Gly Thr Gly 5 Arg Arg Gly Gln Leu His Pro Thr Ala Gly Pro Gly Val His Arg Arg 70 10 Ala Cys Pro Ser Gln Gln Leu Pro His Arg Leu Gly Pro Gly Val Pro 90 Cys Pro Ser Pro Ser Leu Thr Pro Val Leu Pro Ser Trp Thr Gln Ser 105 15 100 Trp Cys Gly Leu Pro Gly Tyr Thr Ser Ser Ser 120 115 20 (2) INFORMATION FOR SEQ ID NO: 631: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 22 amino acids 25 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 631: Val His Gln Leu His Gln Ala Val Gln Gly Cys Ala Leu Gly Arg Pro 30 5 Gly Phe Pro Gly Gly Pro 35 (2) INFORMATION FOR SEQ ID NO: 632: (i) SEQUENCE CHARACTERISTICS: 40 (A) LENGTH: 42 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 632: 45 Pro Thr His Ser Gly His His Lys Ser His Pro Gly Pro Ala Gly Gly Asp Tyr Asn Arg Cys Asp Arg Pro Gly Gln Val His Leu His Asn Pro 50 Arg Gly Thr Gly Arg Arg Gly Gln Leu His 40 55 (2) INFORMATION FOR SEQ ID NO: 633: (i) SEQUENCE CHARACTERISTICS: 60 (A) LENGTH: 55 amino acids

	(B) TYPE: amino acid(D) TOPOLOGY: linear(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 633:
5	Leu His Pro Thr Ala Gly Pro Gly Val His Arg Arg Ala Cys Pro Ser 1 5 10 15
••	Gln Gln Leu Pro His Arg Leu Gly Pro Gly Val Pro Cys Pro Ser Pro 20 25 30
10	Ser Leu Thr Pro Val Leu Pro Ser Trp Thr Gln Ser Trp Cys Gly Leu 35 40 45
15	Pro Gly Tyr Thr Ser Ser Ser 50 55
20	(2) INFORMATION FOR SEQ ID NO: 634:
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 276 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 634:
	Ser Leu Arg Arg Pro Arg Ser Ala Ala Xaa Gln Thr Leu Thr Thr Phe 1 5 10 15
30	Leu Ser Ser Val Ser Ser Ala Ser Ser Ser Ala Leu Pro Gly Ser Arg 20 25 30
25	Glu Pro Cys Asp Pro Arg Ala Pro Pro Pro Pro Arg Ser Gly Ser Ala 35 40 45
35	Ala Ser Cys Cys Ser Cys Cys Cys Ser Cys Pro Arg Arg Ala Pro 50 55 60
40	Leu Arg Ser Pro Arg Gly Ser Lys Arg Arg Ile Arg Gln Arg Glu Val 65 70 75 80
	Val Asp Leu Tyr Asn Gly Met Cys Leu Gln Gly Pro Ala Gly Val Pro 85 90 95
45	Gly Arg Asp Gly Ser Pro Gly Ala Asn Gly Ile Pro Gly Thr Pro Gly 100 105 110
50	Ile Pro Gly Arg Asp Gly Phe Lys Gly Glu Lys Gly Glu Cys Leu Arg 115 120 125
50	Glu Ser Phe Glu Glu Ser Trp Thr Pro Asn Tyr Lys Gln Cys Ser Trp 130 135 140
55	Ser Ser Leu Asn Tyr Gly Ile Asp Leu Gly Lys Ile Ala Glu Cys Thr 145 150 155 160
	Phe Thr Lys Met Arg Ser Asn Ser Ala Leu Arg Val Leu Phe Ser Gly

170

Ser Leu Arg Leu Lys Cys Arg Asn Ala Cys Cys Gln Arg Trp Tyr Phe

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	180	185	190
_	Thr Phe Asn Gly Ala Glu 195	Cys Ser Gly Pro Leu I 200	Pro Ile Glu Ala Ile 205
5	Ile Tyr Leu Asp Gln Gly 210	Ser Pro Glu Met Asn 9	Ser Thr Ile Asn Ile 220
10	His Arg Thr Ser Ser Val 225 230		Gly Ile Gly Ala Gly 240
	Leu Val Asp Val Ala Ile 245	Trp Val Gly Thr Cys	Ser Asp Tyr Pro Lys 255
15	Gly Asp Ala Ser Thr Gly 260	Trp Asn Ser Val Ser 265	Arg Ile Ile Ile Glu 270
20	Glu Leu Pro Lys 275		
	(2) INFORMATION FOR SE	Q ID NO: 635:	
25		ARACTERISTICS: TH: 61 amino acids : amino acid	
30	(D) TOPO (xi) SEQUENCE D	OLOGY: linear ESCRIPTION: SEQ ID NO	
	Ser Leu Arg Arg Pro Ar 1 5	g Ser Ala Ala Xaa Gin 10	The Leu The The File
35	Leu Ser Ser Val Ser Se 20	er Ala Ser Ser Ser Ala 25	Leu Pro Gly Ser Arg 30
	Glu Pro Cys Asp Pro Ax 35	g Ala Pro Pro Pro Pro 40	Arg Ser Gly Ser Ala 45
40	Ala Ser Cys Cys Ser Cy 50	s Cys Cys Ser Cys Pro 55	Arg Arg 60
45	(2) INFORMATION FOR S	EQ ID NO: 636:	
	(A) LEA (B) TYE	HARACTERISTICS: NGTH: 52 amino acids NE: amino acid	
50	(D) TOI (xi) SEQUENCE	OLOGY: linear DESCRIPTION: SEQ ID N	o: 636:
55	Arg Ala Pro Leu Arg S	er Pro Arg Gly Ser Ly	s Arg Arg Ile Arg Gln 15
<i>JJ</i>	Arg Glu Val Val Asp I 20	eu Tyr Asn Gly Met Cy 25	s Leu Gln Gly Pro Ala · 30
60	Gly Val Pro Gly Arg P	sp Gly Ser Pro Gly Al	a Asn Gly Ile Pro Gly 45

```
Thr Pro Gly Ile
          50
5
     (2) INFORMATION FOR SEQ ID NO: 637:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 52 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 637:
     Thr Pro Gly Ile Pro Gly Arg Asp Gly Phe Lys Gly Glu Lys Gly Glu
15
     Cys Leu Arg Glu Ser Phe Glu Glu Ser Trp Thr Pro Asn Tyr Lys Gln
20
      Cys Ser Trp Ser Ser Leu Asn Tyr Gly Ile Asp Leu Gly Lys Ile Ala
               35
      Glu Cys Thr Phe
25
          50
      (2) INFORMATION FOR SEQ ID NO: 638:
30
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 66 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 638:
35
      Phe Thr Lys Met Arg Ser Asn Ser Ala Leu Arg Val Leu Phe Ser Gly
      Ser Leu Arg Leu Lys Cys Arg Asn Ala Cys Cys Gln Arg Trp Tyr Phe
 40
                                        25
       Thr Phe Asn Gly Ala Glu Cys Ser Gly Pro Leu Pro Ile Glu Ala Ile
 45
       Ile Tyr Leu Asp Gln Gly Ser Pro Glu Met Asn Ser Thr Ile Asn Ile
       His Arg
 50
        65
       (2) INFORMATION FOR SEQ ID NO: 639:
 55
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 51 amino acids
```

(B) TYPE: amino acid(D) TOPOLOGY: linear(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 639:

Arg Thr Ser Ser Val Glu Gly Leu Cys Glu Gly Ile Gly Ala Gly Leu Val Asp Val Ala Ile Trp Val Gly Thr Cys Ser Asp Tyr Pro Lys Gly 25 Asp Ala Ser Thr Gly Trp Asn Ser Val Ser Arg Ile Ile Ile Glu Glu 35 10 Leu Pro Lys 50 15 (2) INFORMATION FOR SEQ ID NO: 640: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 26 amino acids 20 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 640: Thr Lys Lys Glu Asn Cys Arg Pro Ala Ser Leu Met Asn Ile Asp Thr 25 Lys Ile Leu Asn Lys Ile Leu Met Asn Gln 20 30 (2) INFORMATION FOR SEQ ID NO: 641: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 214 amino acids 35 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 641: Met Cys Asn Leu Pro Ile Lys Val Val Cys Arg Ala Asn Ala Glu Tyr 40 1 Met Ser Pro Ser Gly Lys Val Pro Xaa Xaa His Val Gly Asn Gln Val 25 45 Val Ser Glu Leu Gly Pro Ile Val Gln Phe Val Lys Ala Lys Gly His Ser Leu Ser Asp Gly Leu Glu Glu Val Gln Lys Ala Glu Met Lys Ala 50 Tyr Met Glu Leu Val Asn Asn Met Leu Leu Thr Ala Glu Leu Tyr Leu 70 Gln Trp Cys Asp Glu Ala Thr Val Gly Xaa Ile Thr His Xaa Arg Tyr 55 90 Gly Ser Pro Tyr Pro Trp Pro Leu Xaa His Ile Leu Ala Tyr Gln Lys 105 110 . 100 60

-	Gln Trp Glu Val Lys Arg Lys Xaa Lys Ala Ile Gly Trp Gly Lys Lys 115 120 125
5	Thr Leu Asp Gln Val Leu Glu Asp Val Asp Gln Cys Cys Gln Ala Leu 130 135 140
	Ser Gln Arg Leu Gly Thr Gln Pro Tyr Phe Phe Asn Lys Gln Pro Thr 145 150 155 160
10	Glu Leu Asp Ala Leu Val Phe Gly His Leu Tyr Thr Ile Leu Thr Thr 165 170 175
15	Gln Leu Thr Asn Asp Glu Leu Ser Glu Lys Val Lys Asn Tyr Ser Asn 180 185 190
	Leu Leu Ala Phe Cys Arg Arg Ile Glu Gln His Tyr Phe Glu Asp Arg 195 200 205
20	Gly Lys Gly Arg Leu Ser 210
25	(2) INFORMATION FOR SEQ ID NO: 642: (i) SEQUENCE CHARACTERISTICS:
30	(A) LENGTH: 44 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 642:
	Met Cys Asn Leu Pro Ile Lys Val Val Cys Arg Ala Asn Ala Glu Tyr 1 5 10 15
35	Met Ser Pro Ser Gly Lys Val Pro Xaa Xaa His Val Gly Asn Gln Val 20 25 30
40	Val Ser Glu Leu Gly Pro Ile Val Gln Phe Val Lys 35 40
-	(2) INFORMATION FOR SEQ ID NO: 643:
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 643: Phe Val Lys Ala Lys Gly His Ser Leu Ser Asp Gly Leu Glu Glu Val
55	Gln Lys Ala Glu Met Lys Ala Tyr Met Glu Leu Val Asn Asn Met Le 20 25 30
	Leu Thr Ala Glu Leu Tyr Leu Gln Trp Cys Asp Glu 35 40

(2) INFORMATION FOR SEQ ID NO: 644:

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
5
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 644:
     Leu Gln Trp Cys Asp Glu Ala Thr Val Gly Xaa Ile Thr His Xaa Arg
10
       1
      Tyr Gly Ser Pro Tyr Pro Trp Pro Leu Xaa His Ile Leu Ala Tyr Gln
15
      Lys Gln Trp Glu Val Lys Arg Lys Kaa Lys Ala Ile Gly Trp Gly Lys
                                   40
      Lys Thr Leu
20
           50
       (2) INFORMATION FOR SEQ ID NO: 645:
25
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 43 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 645:
30
       Asp Gln Val Leu Glu Asp Val Asp Gln Cys Cys Gln Ala Leu Ser Gln
       Arg Leu Gly Thr Gln Pro Tyr Phe Phe Asn Lys Gln Pro Thr Glu Leu
 35
                                        25
       Asp Ala Leu Val Phe Gly His Leu Tyr Thr Ile
                35
 40
        (2) INFORMATION FOR SEQ ID NO: 646:
               (i) SEQUENCE CHARACTERISTICS:
 45
                     (A) LENGTH: 41 amino acids
                     · (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 646:
  50
        Leu Thr Thr Gln Leu Thr Asn Asp Glu Leu Ser Glu Lys Val Lys Asn
                                             10
        Tyr Ser Asn Leu Leu Ala Phe Cys Arg Arg Ile Glu Gln His Tyr Phe
  55
        Glu Asp Arg Gly Lys Gly Arg Leu Ser
```

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(2) INFORMATION FOR SEQ ID NO: 647:
            (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 70 amino acids
5
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 647:
     Met Xaa Xaa Xaa Asn Ser His Ile Thr Ile Fhe Thr Leu Asn Val Asn
10
       1
     Gly Leu Asn Ala Pro Asn Glu Arg His Arg Leu Ala Asn Trp Ile Gln
15
      Ser Gln Asp Gln Val Cys Cys Ile Gln Glu Thr His Leu Thr Gly Arg
                                   40
      Asp Thr His Arg Leu Lys Ile Lys Gly Trp Arg Lys Ile Tyr Gln Ala
20
      Asn Gly Lys Gln Lys Lys
 25
       (2) INFORMATION FOR SEQ ID NO: 648:
              (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 28 amino acids
 30
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 648:
       Phe Thr Leu Asn Val Asn Gly Leu Asn Ala Pro Asn Glu Arg His Arg
 35
                                            10
        Leu Ala Asn Trp Ile Gln Ser Gln Asp Gln Val Cys
                     20
  40
        (2) INFORMATION FOR SEQ ID NO: 649:
                (i) SEQUENCE CHARACTERISTICS:
  45
                       (A) LENGTH: 17 amino acids
                       (B) TYPE: amino acid
                       (D) TOPOLOGY: linear
                (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 649:
   50
         Thr His Leu Thr Gly Arg Asp Thr His Arg Leu Lys Ile Lys Gly Trp
                                              10
         Arg
   55
         (2) INFORMATION FOR SEQ ID NO: 650:
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(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 14 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 650:
 5
     Gly Trp Arg Lys Ile Tyr Gln Ala Asn Gly Lys Gln Lys Lys
10
      (2) INFORMATION FOR SEQ ID NO: 651:
             (i) SEQUENCE CHARACTERISTICS:
15
                     (A) LENGTH: 54 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 651:
      Ile Tyr His Leu His Ser Trp Ile Phe Phe His Phe Lys Arg Ala Phe
20
      Cys Met Cys Phe Ile Thr Met Lys Val Ile His Ala His Cys Ser Lys
                                       25
25
      Leu Arg Lys Cys Xaa Asn Ala Gln Ile Ser Val Phe Cys Thr Thr Leu
               35
      Thr Ala Ser Tyr Pro Thr
30
           50
       (2) INFORMATION FOR SEQ ID NO: 652:
35
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 23 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
40
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 652:
       Ile Tyr His Leu His Ser Trp Ile Phe Phe His Phe Lys Arg Ala Phe
                                            10
         1
                         5
 45
       Cys Met Cys Phe Ile Thr Met
                    20
 50
       (2) INFORMATION FOR SEQ ID NO: 653:
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 31 amino acids
                      (B) TYPE: amino acid
 55
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 653:
       Lys Val Ile His Ala His Cys Ser Lys Leu Arg Lys Cys Xaa Asn Ala
                                            10
 60
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Gln Ile Ser Val Phe Cys Thr Thr Leu Thr Ala Ser Tyr Pro Thr 25 20 5 (2) INFORMATION FOR SEQ ID NO: 654: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 58 amino acids 10 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 654: Trp Asn Leu Leu Trp Tyr Phe Gln Arg Leu Arg Leu Pro Ser Ile Leu 15 5 1 10 Pro Gly Leu Val Leu Ala Ser Cys Asp Gly Pro Ser Xaa Ser Gln Ala 20 Pro Ser Pro Trp Leu Thr Pro Asp Pro Ala Ser Val Gln Val Arg Leu 40 Leu Trp Asp Val Leu Thr Pro Asp Pro Asn 25 (2) INFORMATION FOR SEQ ID NO: 655: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 54 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 655: 35 Gln Arg Gly Ile Tyr Arg Glu Ile Leu Phe Leu Thr Met Ala Ala Leu 10 Gly Lys Asp His Val Asp Ile Val Ala Phe Asp Lys Lys Tyr Lys Ser 40 Ala Phe Asn Lys Leu Ala Ser Ser Met Gly Lys Glu Glu Leu Arg His 35 40 45 Arg Arg Ala Gln Met Pro 50 50 (2) INFORMATION FOR SEQ ID NO: 656: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 656: Trp Asn Leu Leu Trp Tyr Phe Gln Arg Leu Arg Leu Pro Ser Ile Leu 1 5 10

Pro Gly Leu Val Leu Ala Ser 20

5																
	(2) INFORMATION FOR SEQ ID NO: 657:															
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 191 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 657:															
15	Glu 1	Ąsp	Asp	Gly	Phe 5	Asn	Arg	Ser	Ile	His 10	Glu	Val	Ile	Leu	Lys 15	Asn
	Ile	Thr	Trp	Туг 20	Ser	Glu	Arg	Val	Leu 25	Thr	Glu	Ile	Ser	Leu 30	Gly	Ser
20	Leu	Leu	Ile 35	Leu	Val	Val	Ile	Arg 40	Thr	Ile	Gln	Tyr	Asn 45	Met	Thr	Arg
25	Thr	Arg 50	Asp	Lys	Туг	Leu	His 55	Thr	Asn	Суѕ	Leu	Ala 60	Ala	Leu	Ala	Asn
6.J	Met 65	Ser	Ala	Gln	Phe	Arg 70	Ser	Leu	His	Gln	Туг 75	Ala	Ala	Gln	Arg	Ile 80
30	Ile	Ser	Leu	Phe	Ser 85	Leu	Leu	Ser	Lys	Lys 90	His	Asn	Lys	Val	Leu 95	Glu
	Gln	Ala	Thr	Gln 100	Ser	Leu	Arg	Gly	Ser 105		Ser	Ser	Asn	Asp 110	Val	Pro
35	Leu	Pro	Asp 115	-	Ala	Gln	Asp	Leu 120		Val	Ile	Glu	Glu 125	Val	Ile	Arg
40	Met	Met 130	Leu	Glu	Ile	Ile	Asn 135		Cys	Leu	Thr	Asn 140	Ser	Leu	His	His
	Asn 145		Asn	Leu	Val	Туг 150		Leu	Leu	Tyr	Lys 155		Asp	Leu	Phe	Glu 160
45	Gln	Phe	Arg	Thr	His 165		Ser	Phe	Gln	Asp 170		Met	Gln	Asn	Ile 175	
	Leu	Val	Ile	Ser 180		Phe	Ser	Ser	Arg 185		Leu	Gln	Ala	Gly 190		
50																
	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	658:							
55			(i)		(A) 1 (B) 1	E CHA LENG LYPE LOPOI	CH: :	38 ar ino a	mino acid		ls					
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 658:															

 $60\,$ Glu Asp Asp Gly Phe Asn Arg Ser Ile His Glu Val Ile Leu Lys Asn

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	1				5					10					15	
_	Ile	Thr	Trp	Туг 20	Ser	Glu	Arg	Val	Leu 25	Thr	Glu	Ile	Ser	Leu 30	Gly	Ser
5	Leu	Leu	11e 35	Leu	Val	Val										
10	(2)	INFO	ORMA!	rion	FOR	SEQ	ID N	ю: 6	59:							
15				(A) L B) T D) T	ENGT YPE: OPOL	i: 5: ami OGY:	3 am no ac line	ino a cid ear	acid		: 659	9 :			
20	Arg 1	Thr	Ile	Gln	Tyr 5	Asn	Met	Thr	Arg	Thr 10	Arg	Asp	Lys	Tyr	Leu 15	His
	Thr	Asn	Cys	Leu 20	Ala	Ala	Leu	Ala	Asn 25	Met	Ser	Ala	Gln	Phe 30	Arg	Ser
25	Leu	His	Gln 35	Tyr	Ala	Ala	Gln	Arg 40	Ile	Ile	Ser	Leu	Phe 45	Ser	Leu	Leu
30	Ser	Lys 50	Lys	His	Asn											
	(2)	INF	ORMA	TION	FOR	SEQ	ID 1	ю: е	60:							
35			(i)	((A) L	CHAI ENGT YPE: OPOL	H: 5 ami	6 am no a	ino d		s					
40			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 66	0:			
	Ser 1	_	Leu	Thr	Asn 5	Ser	Leu	His	His	Asn 10	Pro	Asn	Leu	Val	Tyr 15	Ala
45	Leu	Leu	Tyr	Lys 20	_	Asp	Leu	Phe	Glu 25	Gln	Phe	Arg	Thr	His 30	Pro	Ser
	Phe	Gln	Asp 35	Ile	Met	Gln	Asn	Ile 40	Asp	Leu	Val	Ile	Ser 45	Phe	Phe	Ser
50	Ser	Arg 50		Leu	Gln	Ala	Gly 55	Ser								
5 5	(2)	INF	ORMA	TION	FOR	SEQ	ID I	NO: (661:							
			(i)	SEQU (CHA ENGI					s					
60						YPE:										

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 661:														
5	Lys Ly 1	s His	Asn	Lys 5	Val	Leu	Glu	Gln	Ala 10	Thr	Gln	Ser	Leu	Arg 15	Gly
3	Ser Le	u Ser	Ser 20	Asn	Asp	Val	Pro	Leu 25	Pro	Asp	Тут	Ala	Gln 30	Asp	
10	(2) IN	IFORMA	TION	FOR	SEQ	ID N	ю: 6	62:							
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 125 amino acids															
15			(B) T D) T	YPE: OPOL	ami OGY:	no a	cid ear							
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 662:															
20	Met Al	la Asp	Ile	Gln 5	Thr	Glu	Arg	Ala	Tyr 10	Gln	Lys	Gln	Pro	Thr 15	Ile
	Phe Gl	ln Asn	Lys 20	Lys	Arg	Val	Leu	Leu 25	Gly	Glu	Thr	Gly	Lys 30	Glu	Lys
25	Leu Pr	o Arg		Thr	Asn	Lys	Asn 40	Ile	Gly	Leu	Gly	Phe 45	Lys	Asp	Thr
30	Pro Ai	rg Arg 50	Leu	Leu	Arg	Gly 55	Thr	Туг	Ile	Asp	Lys 60	Lys	Cys	Pro	Phe
50	Thr G	ly Asn	Val	Ser	Ile 70	Arg	Gly	Arg	Ile	Leu 75	Ser	Gly	Val	Val	Thr 80
35	Gln As	sp Glu	Asp	Ala 85	Glu	Asp	His	Cys	His 90	Pro	Pro	Arg	Leu	Ser 95	Ala
	Leu H	is Pro	Gln 100	Val	Gln	Pro	Leu	Arg 105	Glu	Ala	Pro	Gln	Glu 110	His	Val
40	Cys T	hr Pro		Pro	Leu	Leu	Gln 120	Gly	Arg	Pro	Asp	Arg 125			
• •															
45	(2) I	NFORM			-										
		(1)	SEQU	ENCE (A) I						ls					
50				(B) I (D) I											
		(xi) SEC						EQ I	D NO	: 66	3:			
55	Met L	ys Mei	: Gln	Arg 5		Ile	Val	Ile	Arg 10	_	Asp	Тут	Leu	His 15	
JJ	Ile A	rg Ly:	3 Tyr		Arg	Phe	Glu	Lys 25		His	Lys	Asn	Met 30		Val
60	Kis L	eu Sei 3!		Cys	Phe	Arg	Asp 40		Gln	Ile	Gly	Asp 45		Val	Thr

	Val Gly Slu Cys Arg Pro Les Sar Lys Thr Val Arg Fhe Asm Val Les 50 55 60
5	Lys Val Thr Lys Ala Ala Gly Thr Lys Lys Gln Phe GLL Lys Phe 65 . 70 . 75
10	(2) ENFORMATION FOR SEQ ID NO: 664:
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids (3) TIFE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 684:
20	Met Ala Asp Ile Gln Thr Glu Arg Ala Tyr Gln Lys Glr Pro Thr Ile 1 5 10 13 Phe Gln Asn Lys Lys Arg Val Leu Leu Gly Glu Thr Gly Lys 20 25 21
25	(2) INGTOFMATION FOR SEQ ID NO: 565:
30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 58 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 668:
35	Lys Leu Pro Arg Val Thr Asm Dys Asm Ile Sly Leu Sly Phe Lys Asp 1 13 13
	The Pro Arg Arg Leu Leu Arg Gly The Tyr Tle Asp Lys Lys Tys Sto 20 35 31
40	Phe Thr Gly Asn Val Ser Ile Arg Gly Arg Ile Leu Ser Gly Val Val 35 43 45
45	Thr Gln Asp Glu Asp Ala Glu Asp His Cys 50 55
50	(2) INFORMATION FOR SEQ ID NO: 665:
50	(i) SEQUENCE CHRRACTERISTICS: (A) LENGTH: 38 mains acids
	(E) TYPE: amino acid (C) TOPOLOGY: linear
55	

685

```
Gln Gly Arg Pro Asp Arg
             35
 5
      (2) INFORMATION FOR SEQ ID NO: 667:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 36 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 667:
15
     Met Lys Met Gln Arg Thr Ile Val Ile Arg Arg Asp Tyr Leu His Tyr
      Ile Arg Lys Tyr Asn Arg Phe Glu Lys Arg His Lys Asn Met Ser Val
20
      His Leu Ser Pro
25
      (2) INFORMATION FOR SEQ ID NO: 668:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 43 amino acids
30
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 668:
      Cys Phe Arg Asp Val Gln Ile Gly Asp Ile Val Thr Val Gly Glu Cys
35
      Arg Pro Leu Ser Lys Thr Val Arg Phe Asn Val Leu Lys Val Thr Lys
40
      Ala Ala Gly Thr Lys Lys Gln Phe Gln Lys Phe
               35
45
      (2) INFORMATION FOR SEQ ID NO: 669:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amino acids
                    (B) TYPE: amino acid
50
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 669:
      Pro Arg Arg Leu Leu Arg Gly Thr Tyr Ile Asp Lys Lys Cys Pro Phe
                        5
                                 10
55
      Thr Gly Asn Val Ser Ile Arg Gly Arg Ile Leu Ser Gly Val Val Thr
                   20
                                      25
      Gln
```

```
(2) INFORMATION FOR SEQ ID NO: 670:
 5
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 60 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
10
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 670:
      Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met
       1
                        5
                                           10
15
     Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg
      Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln Arg Ala Lys
                                  40
20
     Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly
          50
                             55
25
      (2) INFORMATION FOR SEQ ID NO: 671:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 67 amino acids
30
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 671:
      Thr Arg Met Ile Asp Leu Leu Glu Glu Tyr Met Val Tyr Arg Lys His
35
      Thr Tyr Xaa Arg Leu Asp Gly Ser Ser Lys Ile Ser Glu Arg Arg Asp
40
     Met Val Ala Asp Phe Gln Asn Arg Asn Asp Ile Phe Val Phe Leu Leu
              35
     Ser Thr Arg Ala Gly Gly Leu Gly Ile Asn Leu Thr Ala Xaa Asp Thr
                              55
45
     Val His Phe
      65
50
      (2) INFORMATION FOR SEQ ID NO: 672:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 32 amino acids
55
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 672:
      Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met
60
                        5
                                          10
```

```
Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg
                                       25
 5
10
      (2) INFORMATION FOR SEQ ID NO: 673:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 31 amino acids
                     (B) TYPE: amino acid
15
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 673:
      Val Tyr Arg Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln
                                           10
20
      Arg Ala Lys Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly
                                       25
25
      (2) INFORMATION FOR SEQ ID NO: 674:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amino acids
30
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 674:
      Thr Arg Met Ile Asp Leu Leu Glu Glu Tyr Met Val Tyr Arg Lys His
35
                                          10
      Thr Tyr Xaa Arg Leu Asp Gly Ser Ser Lys Ile Ser Glu Arg Arg Asp
                   20
                                      25
40
      Met
45
      (2) INFORMATION FOR SEQ ID NO: 675:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 38 amino acids
                     (B) TYPE: amino acid
50
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 675:
      Arg Arg Asp Met Val Ala Asp Phe Gln Asn Arg Asn Asp Ile Phe Val
                                          10
55
      Phe Leu Leu Ser Thr Arg Ala Gly Gly Leu Gly Ile Asn Leu Thr Ala
      Xaa Asp Thr Val His Phe
60
               35
```

5	(2) INFORMATION FOR SEQ ID NO: 676:
3	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 37 amino acids
	(B) TYPE: amino acid
10	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 676:
	Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met 1 5 10 15
15	Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg 20 25 30
	Leu Ile Cys Lys Gly
20	35
	(2) INFORMATION FOR SEQ ID NO: 677:
25	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 37 amino acids (B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 677:
30	
	Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met 1 5 10 15
35	Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg 20 25 30
	Leu Ile Cys Lys Gly 35
40	
	(2) INFORMATION FOR SEQ ID NO: 678:
	(2) INCOMMITON FOR SEQ ID NO. 078.
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 29 amino acids
	(B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 678:
50	
50	Arg Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln Arg Ala 1 5 10 15
	Lys Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly
55	20 25
<i>JJ</i>	
	(2) INFORMATION FOR SEQ ID NO: 679:
60	(i) SEQUENCE CHARACTERISTICS:

	(A) LENGTH: 364 amino acids (B) TYPE: amino acid															
							.OGY:									
			(xi)				SCRI			EO T	ח אור	. 67	۹.			
5			•										٠.			
	Met 1	Ser	Leu	His	Gly 5		Arg	Lys	Glu	Ile 10	Тут	Lys	Tyr	Glu	Ala 15	Pro
10	Trp	Thr	Val	Туг 20	Ala	Met	Asn	Trp	Ser 25	Val	Arg	Pro	Asp	Lys 30	Arg	Phe
	Arg	Leu	Ala 35	Leu	Gly	Ser	Phe	Val 40	Glu	Glu	Туг	Asn	Asn 45	Lys	Val	Gln
15	Leu	Val 50	Gly	Leu	Asp	Glu	Glu 55	Ser	Ser	Glu	Phe	Ile 60	Суз	Arg	Asn	Thr
20	Phe 65	Asp	His	Pro	Tyr	Pro 70	Thr	Thr	Lys	Leu	Met 75	Trp	Ile	Pro	Asp	Thr 80
-0	Lys	Gly	Val		Pro 85	Asp	Leu	Leu	Ala	Thr 90	Ser	Gly	Asp	Tyr	Leu 95	Arg
25	Val	Trp	Arg	Val 100	Gly	Glu	Thr	Glu	Thr 105	Arg	Leu	Glu	Cys	Leu 110	Leu	Asn
	Asn	Asn	Lys 115	Asn	Ser	Asp	Phe	Cys 120	Ala	Pro	Leu	Thr	Ser 125	Phe	Asp	Trp
30	Asn	Glu 130	Val	Asp	Pro	Tyr	Leu 135	Leu	Gly	Thr	Ser	Ser 140	Ile	Asp	Thr	Thr
35	Cys 145	Thr	Ile	Trp	Gly	Leu 150	Glu	Thr	Gly	Gln	Val 155	Leu	Gly	Arg	Val	Asn 160
	Leu	Val	Ser	Gly	His 165	Val	Lys	Thr	Gln	Leu 170	Ile	Ala	His	Asp	Lys 175	Glu
40	Val	Tyr	Asp	Ile 180	Ala	Phe	Ser	Arg	Ala 185	Gly	Gly	Gly	Arg	Asp 190	Met	Phe
	Ala	Ser	Val 195	Gly	Ala	Asp	Gly	Ser 200	Val	Arg	Met	Phe	Asp 205	Leu	Arg	His
45	Leu	Glu 210	His	Ser	Thr	Ile	Ile 215	Tyr	Glu	Asp	Pro	Gln 220	His	His	Pro	Leu
50	Leu 225	Arg	Leu	Cys	Trp	Asn 230	Lys	Gln	Asp	Pro	Asn 235	Tyr	Leu	Ala	Thr	Met 240
	Ala	Met	Asp	Gly	Met 245	Glu	Val	Val	Ile	Leu 250	Asp	Val	Arg	Val	Pro 255	Ala
55	His	Leu	Xaa	Pro 260	Gly	Thr	Thr	Ile	Glu 265	His	Val	Ser	Met	Ala 270	Leu	Leu
	Gly	Pro	His 275	Ile	His	Pro	Ala	Thr 280	Ser	Ala	Leu	Gln	Arg 285	Met	Thr	Thr
60	Arg	Leu	Ser	Ser	Gly	Thr	Ser	Ser	Lys	Cys	Pro	Glu	Pro	Leu	Arg	Thr

•		290					295					300				
5	Leu 305	Ser	Trp	Pro	Thr	Gln 310	Leu	Xaa	Gly	Glu	Ile 315	Asn	Asn	Val	Gln	Trp 320
3	Ala	Ser	Thr	Gln	Pro 325	Glu	Leu	Ser	Pro	Ser 330	Ala	Thr	Thr	Thr	Ala 335	Trp
10	Arg	Tyr	Ser	G1u 340	Cys	Ser	Val	Gly	Gly 345	Ala	Val	Pro.	Thr	Arg 350	Gln	Gly
	Leu	Leu	Туг 355	Phe	Leu	Pro	Leu	Pro 360	His	Pro	Gln	Ser				
15																
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: 6	580:							
20			(i) : (xi)	- (. (.	A) L B) T D) T	engt YPE : OPOL	H: 1 ami OGY:	36 a no a lin	mino cid ear	aci		: 68	D :			
25	Met 1	Ser	Leu	His	Gly 5	Lys	Arg	Lys	Glu	Ile 10	Tyr	Lys	Туг	Glu	Ala 15	Pro
30	Trp	Thr	Val	Туг 20	Ala	Met	Asn	Trp	Ser 25	Val	Arg	Pro	Asp	Lys 30	Arg	Phe
	Arg	Leu	Ala 35	Leu	Gly	Ser	Phe	Val 40	Glu	Glu	Tyr	Asn	Asn 45	Lys	Val	Gln
35	Leu	Val 50	Gly	Leu	Asp	Glu	Glu 55	Ser	Ser	Glu	Phe	Ile 60	Суз	Arg	Asn	Thr
	Phe 65	Asp	His	Pro	Тут	Pro 70	Thr	Thr	Lys	Leu	Met 75	Trp	Ile	Pro	Asp	Thr 80
40	Lys	Gly	Val	Tyr	Pro 85	Asp	Leu	Leu	Ala	Thr 90	Ser	Gly	Asp	Tyr	Leu 95	Arg
45	Val	Trp	Arg	Val 100	Gly	Glu	Thr	Glu	Thr 105	Arg	Leu	Glu	Cys	Leu 110	Leu	Asn
	Asn	Asn	Lys 115	Asn	Ser	Asp	Phe	Cys 120	Ala	Pro	Leu	Thr	Ser 125	Phe	Asp	Trp
50	Asn	Glu 130	Val	Asp	Pro	Tyr	Leu 135	Leu								
55	(2)		ORMAI	SEQUI ()	ENCE A) L B) T	CHAI ENGT YPE:	RACT H: 1	ERIS: 40 au no a	FICS mino cid		ds					
60	-		(xi)					lin PTIO		Юπ	OM C	: 68:	1:			

	Ser 1	Phe	Asp	Trp	Asn 5	Glu	Val	Asp	Pro	Tyr 10	Leu	Leu	Gly	Thr	Ser 15	Ser
5	Ile	Asp	Thr	Thr 20	Cys	Thr	Ile	Trp	Gly 25	Leu	Glu	Thr	Gly	Gln 30	Val	Leu
10	Gly	Arg	Val 35	Asn	Leu	Val	Ser	Gly 40	His	Val	Lys	Thr	Gln 45	Leu	Ile	Ala
	His	Asp 50	Lys	Glu	Val	Tyr	Asp 55	Ile	Ala	Phe	Ser	Arg 60	Ala	Gly	Gly	Gly
15	65			Phe		70					75					80
	Asp	Leu	Arg	His	Leu 85	Glu	His	Ser	Thr	Ile 90	Ile	Tyr	Glu	Asp	Pro 95	Gln
20	His	His	Pro	Leu 100	Leu	Arg	Leu	Cys	Trp 105	Asn	Lys	Gln	Asp	Pro 110	Asn	Tyr
25	Leu	Ala	Thr 115	Met	Ala	Met	Asp	Gly 120	Met	Glu	Val	Val	Ile 125	Leu	Asp	Val
	Arg	Val 130	Pro	Ala	His	Leu	Xaa 135	Pro	Gly	Thr	Thr	Ile 140				
30	(2)	INFO	ORMAT	rion	FOR	SEQ	ID I	NO: 6	582 :							
35				()	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	70 a no a lin	mino cid ear	aci						
			(X1)	SEQ	UENCI	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 68:	2:			
40	Val 1	Gly	Ala	Asp	Gly 5	Ser	Val	Arg	Met	Phe 10	Asp	Leu	Arg	His	Leu 15	Glu
	His	Ser	Thr	Ile 20	Ile	Tyr	Glu	Asp	Pro 25	Gln	His	His	Pro	Leu 30	Leu	Arg
45	Leu	Cys	Trp 35	Asn	Lys	Gln	Asp	Pro 40	Asn	Tyr	Leu	Ala	Thr 45	Met	Ala	Met
50	Asp	Gly 50	Met	Glu	Val	Val	Ile 55	Leu	Asp	Val	Arg	Val 60	Pro	Ala	His	Leu
	Xaa 65	Pro	Gly	Thr	Thr	Ile 70	Glu	His	Val	Ser	Met 75	Ala	Leu	Leu	Gly	Pro 80
55	His	Ile	His	Pro	Ala 85	Thr	Ser	Ala	Leu	Gln 90	Arg	Met	Thr	Thr	Arg 95	Leu
55				Pro Thr 100	85					90					95	

			115					120					125			
5	Thr	Gln 130	Pro	Glu	Leu	Ser	Pro 135	Ser	Ala	Thr	Thr	Thr 140	Ala	Trp	Arg	Tyr
J	Ser 145	Glu	Суз	Ser	Val	Gly 150	Gly	Ala	Val	Pro	Thr 155	Arg	Gln	Gly	Leu	Leu 160
10	Tyr	Phe	Leu	Pro	Leu 165	Pro	His	Pro	Gln	Ser 170						
15	(2)		ORMA:							:						
							H: 2 ami			aci	ds					
20			(xi)				OGY: SCRI			EQ I	D NO	: 68	3:			
	Leu 1	Tyr	Ala	Thr	Ala 5	Thr	Val	Ile	Ser	Ser 10	Pro	Ser	Thr	Glu	Xaa 15	Leu
25	Ser	Gln	Asp	Gln 20	Gly	Asp	Arg	Ala	Ser 25	Leu	Asp	Ala	Ala	Asp 30	Ser	Gly
30	Arg	Gly	Ser 35	Trp	Thr	Ser	Cys	Ser 40	Ser	Gly	Ser	His	Asp 45	Asn	Ile	Gln
	Thr	Ile 50	Gln	His	Gln	Arg	Ser 55	Trp	Glu	Thr	Leu	Pro 60	Phe	Gly	His	Thr
35	His 65	Phe	Asp	Tyr	Ser	Gly 70	Asp	Pro	Ala	Gly	Leu 75	Trp	Ala	Ser	Ser	Ser 80
	His	Met	Asp	Gln	Ile 85	Met	Phe	Ser	Asp	His 90	Ser	Thr	Lys	Tyr	Asn 95	Arg
40	Gln	Asn	Gln	Ser 100	Arg	Glu	Ser	Leu	Glu 105	Gln	Ala	Gln	Ser	Arg 110	Ala	Ser
45	Trp	Ala	Ser 115	Ser	Thr	Gly	Tyr	Trp 120	Gly	Glu	Asp	Ser	Glu 125	Gly	Asp	Thr
		130	Ile				135					140				
50	145		Ser			150					155					160
			Ala		165					170					175	
5 5			Lys	180					185					190		
60	Tyr	Ile	Gly 195	Ile	Pro	Ile	Thr	Asp 200	Phe	Pro	Glu	Gly	His 205	Ser	His	Pro

•	Ala	Arg 210	Lys	Pro	Pro	Asp	Туг 215	Asn	Val	Ala	Leu	Gln 220	Arg	Ser	Arg	Met
5	Val 225	Ala	Arg	Ser	Ser	Asp 230	Thr	Ala	Gly	Pro	Ser 235	Ser	Val	Gln	Gln	Pro 240
	His	Gly	His	Pro	Thr 245	Ser	Ser	Arg	Pro	Val 250	Asn	Lys	Pro	Gln	Trp 255	His
10	Lys	Xaa	Asn	Glu 260	Ser	Asp	Pro	Arg	Leu 265	Ala	Pro	Tyr	Gln	Ser 270	Gln	Gly
15	Phe	Ser	Thr 275	Glu	Glu	Asp	Glu	Asp 280	Glu	Gln	Val	Ser	·Ala 285	Val		
	(2)	INF	ORMA'	TION	FOR	SEQ	ID I	NO: (684:							
20				(A) L B) T D) T	ENGT YPE : OPOL	H: 4 ami OGY:	2 am no a lin	ino cid ear	acid		: 68	4:			
25	His 1		Asp	Gln	Ile 5	Met	Phe	Ser	Asp	His 10	Ser	Thr	Lys	Tyr	Asn 15	Arg
30	Gln	Asn	Gln	Ser 20	Arg	Glu	Ser	Leu	Glu 25	Gln	Ala	Gln	Ser	Arg 30	Ala	Ser
	Trp	Ala	Ser 35	Ser	Thr	Gly	Tyr	Trp 40	_	Glu				7		
35																
	(2)	INF		TION SEQU		_	•			:						
40			(xi)	((A) I (B) I (D) I	YPE:	ami OGY :	no a	cid ear			: 6 8	5:			
45	Ser 1		Thr	Thr	Glu 5		Thr	Lys	Pro	Val		Met	Pro	Ala	His 15	Ile
50	Ala	Val	Ala	Ser 20		Thr	Thr	Lys	Gly 25		Ile	Ala	Arg	Lys 30		Gly
	Arg	Туг	Arg		Pro	Pro	Pro	Thr 40		Pro	Gly	Тух	11e 45	-	Ile	Pro
55	Ile	Thr 50	_													
60	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	686 :							•

	(A) LENGTH: 57 amino acids (B) TYPE: amino acid
5	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 686:
	Val Ala Leu Glm Arg Ser Arg Met Val Ala Arg Ser Ser Asp Thr Ala 1 5 10 15
10	Gly Pro Ser Ser Val Glm Glm Pro His Gly His Pro Thr Ser Ser Arg 20 25 30
15	Pro Val Asn Lys Pro Gln Trp His Lys Xaa Asn Glu Ser Asp Pro Arg 35 40 45
13	Leu Ala Pro Tyr Gln Ser Gln Gly Phe 50 55
20	(2) DIFORMATION FOR SEQ ID NO: 687:
25	(i) SEQUENCE CHARACTERISTICS: (A) LEWSTH: 41 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 687:
30	Cys Leu Leu Phe Val Phe Val Ser Leu Gly Met Arg Cys Leu Phe Trp 1 5 10 15
	Thr Ile Val Tyr Asm Val Leu Tyr Leu Lys His Lys Cys Asm Thr Val 20 25 30
35	Leu Leu Cys Tyr His Leu Cys Ser Ile 35 40
40	(2) INFORMATION FOR SEQ ID NC: 688:
45	(i) SEQUENCE CHERACTERISTICS: (A) LENGTH: 67 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 688:
	Ala Cys Ser Lys Leu Ile Pro Ala Phe Glu Met Val Met Arg Ala Lys
50	1 5 10 15 Asp Asn Val Tyr His Leu Asp Cys Phe Ala Cys Gln Leu Cys Asn Gln 20 25 30
55	Arg Xaa Cys Val Gly Asp Lys Phe Phe Leu Lys Asn Asn Xaa Xaa Leu 35 40 45
	Cys Gln Thr Asp Tyr Glu Glu Gly Leu Met Lys Glu Gly Tyr Ala Pro 50 55 60
60	Xaa Val Arg

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5	(2) INFORMATION FOR SEQ ID NO: 689:
10	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 45 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 689:
	Ser Ala Leu Ser Glu Pro Gly Ala Pro Asp Arg Arg Pro Cys Pro
	1 5 10 15
15	
	Glu Ser Val Pro Arg Arg Pro Asp Asp Glu Gln Trp Pro Pro Pro Thr 20 25 30
	Ala Leu Cys Leu Asp Val Ala Pro Leu Pro Pro Ser Ser
20	35 40 45
25	(2) INFORMATION FOR SEQ ID NO: 690:
23	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 43 amino acids
	(B) TYPE: amino acid
30	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 690:
	,,,,,,
	Pro Val Gly Tyr Leu Asp Lys Gln Val Pro Asp Thr Ser Val Gln Glu 1 5 10 15
	1 5 10 13
35	Thr Asp Arg Ile Leu Val Glu Lys Arg Cys Trp Asp Ile Ala Leu Gly
	20 25 30
	Pro Leu Lys Gln Ile Pro Met Asn Leu Phe Ile
40	35 40
40	
	(2) INFORMATION FOR SEQ ID NO: 691:
45	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 214 amino acids
	(B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 691:
50	(AL) Signatus sessiterium. Seg Is no. USI.
	Ala His Ala Ser Glu Ser Gly Glu Arg Trp Trp Ala Cys Cys Gly Val
	1 5 10 15
	Arg Phe Gly Leu Arg Ser Ile Glu Ala Ile Gly Arg Ser Cys Cys His
55	20 25 30
	Asp Gly Pro Gly Gly Leu Val Ala Asn Arg Gly Arg Arg Phe Lys Trp
	35 40 45
6 0	
60	Ala Ile Glu Leu Ser Gly Pro Gly Gly Gly Ser Arg Gly Arg Ser Asp

		50					55					60				
5	Arg 65	Gly	Ser	Gly	Gln	Gly 70	Asp	Ser	Leu	Tyr	Pro 75	Val	Gly	Туг	Leu	Asp 08
J	Lys	Gln	Val	Pro	Asp 85	Thr	Ser	Val	Gln	Glu 90	Thr	Asp	Arg	Ile	Leu 95	Val
10	Glu	Lys	Arg	Суз 100	Trp	Asp	Ile	Ala	Leu 105	Gly	Pro	Leu	Lys	Gln 110	Ile	Pro
	Met	Asn	Leu 115	Phe	Ile	Met	Tyr	Met 120	Ala	Gly	Asn	Thr	Ile 125	Ser	Ile	Phe
15	Pro	Thr 130	Met	Met	Val	Cys	Met 135	Met	Ala	Trp	Arg	Pro 140	Ile	Gln	Ala	Leu
20	Met 145	Ala	Ile	Ser	Ala	Thr 150	Phe	Lys	Met	Leu	Glu 155	Ser	Ser	Ser	Gln	Lys 160
	Phe	Leu	Gln	Gly	Leu 165	Val	Тут	Leu	Ile	Gly 170	Asn	Leu	Met	Gly	Leu 175	Ala
25	Leu	Ala	Val	Туг 180	Lys	Cys	Gln	Ser	Met 185	Gly	Leu	Leu	Pro	Thr 190	His	Ala
	Ser	Asp	Trp 195	Leu	Ala	Phe	Ile	Glu 200	Pro	Pro	Glu	Arg	Met 205	Glu	Phe	Ser
30	Gly	Gly 210	Gly	Leu	Leu	Leu										
35	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	692 :							
40				((A) I (B) I (D) I	ENGI YPE : YOPOI	H: 4 ami OGY:	ino a lino a	ino cid ear	acid): 6 9	2:			
	Ala		Phe	Lys	Met 5		Glu	Ser	Ser	Ser 10		Lys	Phe	Leu	Gln 15	Gly
45	Leu	Val	. Туг	Leu 20		Gly	Asn	Leu	Met 25		Leu	Ala	Leu	Ala 30		Tyr
50	Lys	Cys	Gln 35	Ser	Met	Gly	Leu	Leu 40		Thr	His	Ala	Ser 45		•	
55	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	693 :							
-			(i)		(A) 1 (B) 1	LENG IYPE	rh:	43 ar ino a	nino acid		ds					
60			(xi	SEX		TOPOI				TEXT 1	m Na)· 69	93:			

WO 98/54963

697

PCT/US98/11422

	Pro 1	Val	Gly	Tyr	Leu 5	Asp	Lys	Gln	Val	Pro 10	Asp	Thr	Ser	Val	Gln 15	Glu
5	Thr	Asp	Arg	Ile 20	Leu	Val	Glu	Lys	Arg 25	Cys	Trp	Asp	Ile	Ala 30	Leu	Gly
10	Pro	Leu	Lys 35	Gln	Ile	Pro	Met	Asn 40	Leu	Phe	Ile					
15	(2)	INF	ORMAT	SEQUI () (ENCE A) L B) T D) T	CHA ENGT YPE: OPOL	RACT H: 4 ami OGY:	ERIS 8 am no a lin	TICS ino cid ear	acid		. 69	1.			
20	Pro 1	Thr	(xi) Thr											Ile	Gln 15	Ile
25	Arg	Phe	Pro	Ser 20	Phe	Tyr	His	Lys	Leu 25	Val	Asp	Ser	Gly	Arg 30	Met	Arg
	Ser	Lys	Arg 35	Glu	Thr	Arg	Arg	Glu 40	Asp	Ser	Asp	Thr	Lys 45	His	Asn	Leu
30																
35	(2)	INF	ORMA'	rion	FOR	SEQ	ID	NO:	695 :							
40				(A) I B) T D) T	ENGI YPE : OPOI	H: 1 ami OGY:	.67 a .no a : lin	mino cid ear	aci		: 69	5:			
45	Thr 1		His	Ile	Ile 5		Val	Met	Ile	Thr 10		Leu	Arg	Gly	Lys 15	Asp
	Ile	Leu	Ser	Тух 20		Glu	Lys		Ile 25		Val	Gln	Met	Thr 30	Ile	Ala
50	Val	Gly	Thr 35	_	Met	Pro	Pro	Lys 40		Phe	Ser	Arg	Gly 45	Ser	Leu	Val
		50					55					60				
55	65					70					75					Asp 80
60	Arg	Asn	Gln	Arg	Arg 85		Gly	' Asp	Ala	Ala 90	-	Lys	Ala	Ile	Ser 95	_

	Leu	Thr	Thr	Arg 100	Thr	Val	Lys	Lys	Gly 105	Asp	Lys	Glu	Thr	Asp 110	Pro	Asp
5	Phe	Asp	His 115	Суз	Ala	Val	Cys	Ile 120	Glu	Ser	Tyr	Lys	Gln 125	Asn	Asp	Val
	Val	Arg 130	Ile	Leu	Pro	Cys	Lys 135	His	Val	Phe	His	Lys 140	Ser	Cys	Val	Asp
10	Pro 145	Trp	Leu	Ser	Glu	His 150	Cys	Thr	Суз	Pro	Met 155	Суз	Lys	Leu	Asn	Ile 160
15	Leu	Lys	Ala	Leu	Gly 165	Ile	Val									
'no	(2)	INF	ORMA													
20			(1)	(engt Ype:	H: 2 ami	76 a no a			ds					
25				_					N: S	_				-1-		6 3
	Met 1		His	Pro	G1y 5	Thr	Glu	His	Ile	11e	Ala	Val	Met	IIe	Thr 15	Glu
30	Leu	Arg	Gly	Lys 20	Asp	Ile	Leu	Ser	Tyr 25	Leu	Glu	Lys	Asn	Ile 30	Ser	Val
	Gln	. Met	Thr 35		Ala	Val	Gly	Thr 40		Met	Pro	Pro	Lys 45	Asn	Phe	Ser
35	Arg	Gl ₃ 50	/ Ser	Leu	Val	Phe	Val 55	Ser	Ile	Ser	Phe	Ile 60	Val	Leu	Met	Ile
40	Ile 65		: Ser	Ala	Trp	Leu 70	Ile	Phe	Tyr	Phe	Ile 75		Lys	Ile	Arg	Tyr 80
	Thr	: Ası	n Ala	Arg	Asp 85	Arg	Asn	Gln	Arg	Arg 90		Gly	Asp	Ala	Ala 95	
45	Lys	: Ala	a Ile	Ser 100	-	Leu	Thr	Thr	Arg 105	Thr	Val	Lys	Lys	Gly 110		Lys
	Glu	Th	r Asp 115		Asp	Phe	Asp	His 120		Ala	Val	Cys	11e		Ser	Tyr
50	Lys	130		a Asp	Val	Val	Arg 135		: Leu	Pro	Суз	Lys 140		Val	Phe	His
55	Lys 145		r Cys	Val	Asp	Pro 150	-	Leu	Ser	Glu	His 155	-	Thr	Суз	Pro	Met 160
	Cys	. Ly:	s Leu	ı Asn	11e		Lys	: Ala	Leu	Gly 170		Val	. Pro	Asn	175	Pro
60	Cys	5 Th	r Ası	Asn 180		Ala	Phe	. Asp	Met 185		Arg	Lev	Thi	Arg		Gln

	Ala	Val	Asn 195	Arg	Arg	Ser	Ala	Leu 200	Gly	Asp	Leu	Ala	Gly 205	Asp	Asn	Ser
5	Leu	Gly 210	Leu	Glu	Pro	Leu	Arg 215	Thr	Ser	Gly	Ile	Ser 220	Pro	Leu	Pro	Gln
10	Asp 225	Gly	Glu	Leu	Thr	Pro 230	Arg	Thr	Gly	Glu	Ile 235	Asn	Ile	Ala	Val	Thr 240
10	Lys	Glu	Trp	Phe	Ile 245	Ile	Ala	Ser	Phe	Gly 250	Leu	Leu	Ser	Ala	Leu 255	Thr
15	Leu	Cys	Tyr	Met 260		Ile	Arg	Ala	Thr 265		Ser	Leu	Asn	Ala 270	Asn	Glu
	Val	Glu	Trp 275	Phe												
20	(2)	INF	ORMA	TION	FOR	SEO	ID	NO:	697 :							
25	(2)	1141	(i)	SEQU	ENCE (A) I (B) I	CHA LENGI TYPE:	RACT H: (: am:	TERIS 69 ar ino a : lir	TICS mino acid near	ació): 6 9	97 :			
30	Thr 1		. His	; Ile	: Ile		. Val	l Met	: Ile	Thr 10		Leu	Arg	Gly	Lys 19	Asp
35	Ile	e Leu	sez	Tyr 20		Glu	Lys	s Ası	1 Ile 25		Val	Glr	Met	Thr 30		a Ala
	Val	Gly	7 Thi 35		j Met	Pro	Pro	Lys		. Phe	e Ser	: Arg	Gly 49		: Let	ı Val
40	Phe	• Va]		r Ile	e Sex	Phe	11e 59		l Le	ı Met	: Ile	e Ile 60		c Ser	: Ala	a Trp
	Let 69		e Phe	е Тул	r Phe	•										
45	12)	eo pw	ATIO!	vi EOi	P CF	תז ר	NO.	698							
	(2,	, 111/		SEQ												
50			(xi	.) SE	(B) (D)	TYPE TOPO	LOGY	58 a ino (: li IPTI	acid near	:		ro: 6	98:	-		
5 5		r Il 1	e Se	r Ph		e Va 5	l Le	u Me	t Il	e Il 1		r Se	r Al	a Tr		u Ile 5
	Ph	е Ту	r Ph	e Il 2		n Ly	s Il	.e Ar	g Ту 2		r As	n Al	a Ar		p Ar O	g Asn
ራ ስ																

	Gln	Arg	Arg 35	Leu	Gly	Asp	Ala	Ala 40	Lys	Lys	Ala	Ile	Ser 45	Lys	Leu	Thr
5	Thr	Arg 50	Thr	Val	Lys	Lys	Gly 55	Asp	Lys	Glu						
10	(2)			rion Sequ						_						
. ~				(A) L B) T D) T	engt YPE : OPOL	H: 6 ami OGY:	6 am no a lin	ino cid ear	acid						
15			(xi)	SEQ	UENC:	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 69	9:			
	Val 1	Lys	Lys	Gly	Asp 5	Lys	Glu	Thr	Asp	Pro 10	Asp	Phe	Asp	His	Cys 15	Ala
20	Val	Cys	Ile	Glu 20	Ser	Tyr	Lys	Gln	Asn 25	Asp	Val	Val	Arg	Ile 30	Leu	Pro
25	Cys	Lys	His 35	Val	Phe	His	Lys	Ser 40	Cys	Val	Asp	Pro	Trp 45	Leu	Ser	Glu
	His	Суs 50	Thr	Cys	Pro	Met	Cys 55	Lys	Leu	Asn	Ile	Leu 60	Lys	Ala	Leu	Gly
30	Ile 65	Val												ē		
35	(2)	INF		TION SEQU												
			,	(A) L B) T D) T	ENGT YPE:	H: 1 ami	06 a no a	mino cid		ds					
40			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 70	0:			
	Met 1	Thr	His	Pro	Gly 5	Thr	Glu	His	Ile	Ile 10	Ala	Val	Met	Ile	Thr 15	Glu
45	Leu	Arg	Gly	Lys 20	Asp	Ile	Leu	Ser	Туг 25	Leu	Glu	Lys	Asn	Ile 30	Ser	Val
50	Gln	Met	Thr 35	Ile	Ala	Val	Gly	Thr 40	Arg	Met	Pro	Pro	Lys 45	Asn	Phe	Ser
J	Arg	Gly 50	Ser	Leu	Val	Phe	Val 55	Ser	Ile	Ser	Phe	Ile 60	Val	Leu	Met	Ile
55	Ile 65	Ser	Ser	Ala	Trp	Leu 70	Ile	Phe	Tyr	Phe	Ile 75	Gln	Lys	Ile	Arg	Туг 80
	Thr	Asn	Ala	Arg	Asp 85	Arg	Asn	Gln	Arg	Arg 90	Leu	Gly	Asp	Ala	Ala 95	Lys
50	Lys	Ala	Ile	Ser	Lys	Leu	Thr	Thr	Arg	Thr						

5	(2) INFORMATION FOR SEQ ID NO: 701:
10	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 84 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 701:
15	Ala Ala Lys Lys Ala Ile Ser Lys Leu Thr Thr Arg Thr Val Lys Lys 1 5 10 15
	Gly Asp Lys Glu Thr Asp Pro Asp Phe Asp His Cys Ala Val Cys Ile 20 25 30
20	Glu Ser Tyr Lys Gln Asn Asp Val Val Arg Ile Leu Pro Cys Lys His 35 40 45
	Val Phe His Lys Ser Cys Val Asp Pro Trp Leu Ser Glu His Cys Thr 50 55 60
25	Cys Pro Met Cys Lys Leu Asn Ile Leu Lys Ala Leu Gly Ile Val Pro 65 70 75 80
30	Asn Leu Pro Cys
	(2) INFORMATION FOR SEQ ID NO: 702:
35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 86 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 702:
• .	Thr Gln Ala Val Asn Arg Arg Ser Ala Leu Gly Asp Leu Ala Gly Asp 1 5 10 15
45	Asn Ser Leu Gly Leu Glu Pro Leu Arg Thr Ser Gly Ile Ser Pro Leu 20 25 30
	Pro Gln Asp Gly Glu Leu Thr Pro Arg Thr Gly Glu Ile Asn Ile Ala 35 40 45
50	Val Thr Lys Glu Trp Phe Ile Ile Ala Ser Phe Gly Leu Leu Ser Ala 50 55 60
55	Leu Thr Leu Cys Tyr Met Ile Ile Arg Ala Thr Ala Ser Leu Asn Ala 65 70 75 80

Asn Glu Val Glu Trp Phe

	-,		· · · · ·			~~~			•••							
5			(i) E	() ()	A) Li 2) Ti 0) Ti	eigt: Yfe: Opclu	e: 3. a ni: XGY:	41 ar no ac line	nins sid sar	acis		: 79	3:			
10	Pro 1	Leu	His	Gly	7al 5	λia	Asp	#is	Leu	31y 10	Çys	æş,	Pro	Gln	Thr 15	Arg
	Fite	Fhe	Val	Pro 20	320	Asn	Ile	_'/s	3 <u>1-</u> 25	ŢŢŢ	Ile	λia	Leu	Leu 30	Gln	Arg
15	Bly	Ast.	35 35	The	Phe	Lys	Glu	Lys 40	Ile	Ser	λzg	Ala	Ala 45	Phe	His	Asn
20	λia	Val 50	Ala	Val	Val	Ile	Tyr 55	Asn	Asn	Lys	Ser	Lys 50	Glu	Glu	Pro	Val
20	Thr 65	Υετ	ጥ느	His	220	G <u>l</u> y 70	Thr	Glu	His	Ile	Ile 75	Ala	Val	Mec	Ile	Thr 80
25	Glu	Leu	Arg	Gly	Lys 25	ćsy	Ile	Leu	Ser	בעב פעב	Leu	Gla	Lys	Asn	Ile 95	Ser
	Wal	3lm	. Xet	Thr 100	Ile	Ala	Val	зіу	Thr 105	Yrş	Xet	P20	Pro	Lys 110	Asn	Phe
30	Ser	Arg	Gly 115	Ser	Leu	Val	Phe	∵al 120	Ser	Ile	Ser	Phe	Ile 125	٧al	Leu	Met
35	lle	11s	:Ser	Ser	Ala	TTP	Leu 135	Ile	Phe	፲/፲	Phe	11e 140		Lys	Ile	Arg
J J	7yr 145		: Ast	Ala	Arg	750 150		Asn	Gln	Yrg	Arg 155	Leu	Gly	qzA	Ala	Ala 160
40	Lys	Lys	Ale	Ile	Ser 185	-	Leu	<u>Thr</u>	Thr	Arg 170	Thr	Val	Lys	Lys	Gly 175	
	Lys	31:	: The	Asp 180		Ąsp	Phe	Asp	His 185		Ala	Val	Cys	Ile 190		Ser
45	בויי: הייי	Lys	GL- 195	Asn	Asp	Val	Val	æg 200	Ile	Leu	220	Cys	Lys 205	His	Val	Phe
50	His	Ly: 219	s Se≃)	Cys	Val	Asp	Pro 215	_	Leu	Ser	Glu	⊞is 220	_	Thr	Cys	Pro
50	<u>Yet</u> 225	-	s Lys	Leu	. Asn	Ile 230		Lys	Ala	Leu	Gly 235		Val	Pro	Asn	Leu 240
55	Pro	Cy:	Th <u>r</u>	Asp	Asn 245		Ala	?he	Asp	Met 250		Arg	Leu	Thr	Arg 255	
	Slr	: Al	a Val	As:: 260	_) Arg	Ser	Ala	Leu 265	_	ودن	Leu	Ala	. Gly 270	•	Asn
60	201	. 1 .		Tax	. 61.	S	Ton		. The		٠		Sar	· Dro	ופו	Dro

		275		280			285	
5	Gln Asp 290	Gly Glu	Leu Thr	Pro Arg 295	Thr Gly	Glu Ile 300	Asn Ile	Ala Val
J	Thr Lys 305	Glu Trp	Phe Ile 310	Ile Ala	Ser Phe	Gly Leu 315	Leu Ser	Ala Leu 320
10	Thr Leu	Cys Tyr	Met Ile 325	Ile Arg	Ala Thr 3	Ala Ser	Leu Asn	Ala Asn 335
	Glu Val	Glu Trp 340	Phe					
15								
	(2) INF	ORMATION	FOR SEQ	ID NO:	704:			
20		(B) TYPE: D) TOPOL	H: 60 am amino a OGY: lin	ino acids cid		4 :	
25	His Gly	Val Ala	Asp His 5	Leu Gly	Cys Asp 10	Pro Gln	Thr Arg	Phe Phe
30	Val Pro	Pro Asn 20	Ile Lys	Gln Trp	Ile Ala 25	Leu Leu	Gln Arg 30	Gly Asn
30	Cys Thr	Phe Lys 35	Glu Lys	Ile Ser 40	Arg Ala	Ala Phe	His Asn 45	Ala Val
35	Ala Val		Tyr Asn	Asn Lys 55	Ser Lys	Glu Glu 60		
40	(2) INF	ORMATION	FOR SEQ					
		_ ((A) LENGT	TH: 314 a	mino acid	is		
45		((B) TYPE: (D) TOPOI (UENCE DE	OGY: lir		NO: 70	5:	,
	Met Ser 1	Gly Gln	Gly Leu 5	Ala Gly	Phe Phe	Ala Ser	Val Ala	Met Ile 15
50	Cys Ala	lle Ala 20	_	Ser Glu	Leu Ser 25	Glu Ser	Ala Phe	Gly Tyr
55	Phe Ile	Thr Ala	Cys Ala	Val Ile	Ile Leu	Thr Ile	Ile Cys 45	Tyr Leu
<i></i>	Gly Let 50		Leu Glu	Phe Tyr 55	Arg Tyr	Tyr Gln 60		Lýs Leu
60	Glu Gly 65	/ Pro Gly	Glu Gln 70		Lys Leu	Asp Leu 75	Ile Ser	Lys Gly 80

	Glu	Glu	Pro	Arg	Ala 85	Gly	Lys	Glu	Glu	Ser 90	Gly	Val	Ser	Väl	Ser 95	Asn
5	Ser	Gln	Pro	Thr 100	Asn	Glu	Ser	His	Ser 105	Ile	Lys	Ala	Ile	Leu 110	Lys	Asn
10	Ile	Ser	Val 115	Leu	Ala	Phe	Ser	Val 120	Cys	Phe	Ile	Phe	Thr 125	Ile	Thr	Ile
		130		Pro			135					140				_
15	Ser 145	Ser	Thr	Trp	Glu	Arg 150	Tyr	Phe	Ile	Pro	Val 155	Ser	Cys	Phe	Leu	Thr 160
	Phe	Asn	Ile	Phe	Asp 165	Trp	Leu	Gly	Arg	Ser 170	Leu	Thr	Ala	Val	Phe 175	Met
20	Trp	Pro	Gly	Lys 180	Asp	Ser	Arg	Trp	Leu 185	Pro	Ser	Trp	Xaa	Leu 190	Ala	Arg
25	Leu	Val	Phe 195	Val	Pro	Leu	Leu	Leu 200	Leu	Cys	Asn	Ile	Lys 205	Pro	Arg	Arg
	Tyr	Leu 210	Thr	Val	Val	Phe	Glu 215	His	Asp	Ala	Trp	Phe 220	Ile	Phe	Phe	Met
30	Ala 225	Ala	Phe	Ala	Phe	Ser 230	Asn	Gly	Tyr	Leu	Ala 235	Ser	Leu	Cys	Met	Cys 240
	Phe	Gly	Pro	Lys	Lys 245	Val	Lys	Pro	Ala	Glu 250	Ala	Glu	Thr	Ala	Glu 255	Pro
35	Ser	Trp	Pro	Ser 260	Ser	Cys	Val	Trp	Val 265	Trp	His	Trp	Gly	Leu 270	Phe	Ser
1 0	Pro	Ser	Cys 275	Ser	Gly	Gln	Leu	Cys 280	Asp	Lys	Gly	Trp	Thr 285	Glu	Gly	Leu
	Pro	Ala 290	Ser	Leu	Pro	Val	Cys 295	Leu	Leu	Pro	Leu	Pro 300	Ser	Ala	Arg	Gly
1 5	Asp 305	Pro	Glu	Trp	Ser	Gly 310	Gly	Phe	Phe	Phe						
	(2)	INFO	ORMA!	MOI	FOR	SEO	ID N	10: 7	706:							
50				SEQUE	ENCE		RACTI	RIST	rics		đa					
55			(xi)	(1	B) T	YPE: OPOL	ami CGY:	no ao lino	cid ear			. 704	٤.			
	Met 1			Gln						_				Ala	Met 15	Ile
50		Ala	Ile	Ala	-	Gly	Ser	Glu	Leu		Glu	Ser	Ala	Phe		Tyr

•				20					25					30		
5	Phe	Ile	Thr 35	Ala	Cys	Ala	Val	Ile 40	Ile	Leu	Thr	Ile	Ile 45	Суз	Туг	Leu
J	Gly	Leu 50	Pro	Arg	Leu	Glu	Phe 55	Tyr	Arg	Tyr	Tyr	Gln 60	Gln	Leu	Lys	Leu
10	Glu 65	Gly	Pro	Gly	Glu	Gln 70	Glu	Thr	Lys	Leu	Asp 75	Leu	Ile	Ser	Lys	80 Gly
	Glu	Glu	Pro	Arg	Ala 85	Gly	Lys	Glu	Glu	Ser 90	Gly	Val	Ser	Val	Ser 95	Asn
15	Ser	Gln	Pro	Thr 100	Asn	Glu	Ser	His	Ser 105	Ile						
20	(2)	INFO	ORMA!	rion	FOR	SEQ	ID I	vo: 7	707 :							
			(i)	SEQUI	ENCE A) L						•					
25			(xi)	(B) T D) T	YPE: OPOL	ami OGY:	no a lin	cid ear		-	: 70°	7:			
30	Ser 1	Gly	Val	Ser	Val 5	Ser	Asn	Ser	Gln	Pro 10	Thr	Asn	Glu	Ser	His 15	Ser
30	Ile	Lys	Ala	Ile 20	Leu	Lys	Asn	Ile	Ser 25	Val	Leu	Ala	Phe	Ser 30	Val	Cys
35	Phe	Ile	Phe 35	Thr	Ile	Thr	Ile	Gly 40	Met	Phe	Pro	Ala	Val 45	Thr	Val	Glu
	Val	Lys 50	Ser	Ser	Ile	Ala	Gly 55	Ser	Ser	Thr	Trp	Glu 60	Arg	Tyr	Phe	Ile
40	Pro 65	Val	Ser	Cys	Phe	Leu 70	Thr	Phe	Asn	Ile	Phe 75	Asp	Trp	Leu	Gly	Arg 80
	Ser															
45																
	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: 1	708:							
50 .			(i)	(A) L B) T	engt Ype :	H: 9 ami	2 am no a	ino cid		s					
55			(xi)	SEQ	D) T UENC					EQ I	D NO	: 70	8:			
JJ	Thr 1		Gly	Met	Phe 5	Pro	Ala	Val	Thr	Val 10	Glu	Val	Lys	Ser	Ser 15	Ile
60	Ala	Gly	Ser	Ser 20	Thr	Trp	Glu	Arg	Tyr 25	Phe	Ile	Pro	Val	Ser 30	Суз	Phe

	Leu	Thr	35	ASII	TTG	Pne	ASP	40	rea	GIÀ	Arg	ser	45	THE	Ala	vai
5	Phe	Met 50	Trp	Pro	Gly	Lys	Asp 55	Ser	Arg	Trp	Leu	Pro 60	Ser	Trp	Xaa	Leu
10	Ala 65	Arg	Leu	Val	Phe	Val 70	Pro	Leu	Leu	Leu	Leu 75	Cys	Asn	Ile	Lys	Pro 80
	Arg	Arg	Tyr	Leu	Thr 85	Val	Val	Phe	Glu	His 90	Asp	Ala				
15	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	VO: 7	709:							
20				(A) L B) T D) T	engi YPE: OPOL	H: 7 ami OGY:	4 am no a lin	ino a cid ear	acid		: 70:	9:			
25	Phe 1	Gly	Pro	Lys	Lys 5	Val	Lys	Pro	Ala	Glu 10	Ala	Glu	Thr	Ala	Glu 15	Pro
	Ser	Trp	Pro	Ser 20	Ser	Cys	Val	Trp	Val 25	Trp	His	Trp	Gly	Leu 30	Phe	Ser
30	Pro	Ser	Суs 35	Ser	Gly	Gln	Leu	Cys 40	Asp	Lys	Gly	Trp	Thr 45	Glu	Gly	Leu
35	Pro	Ala 50	Ser	Leu	Pro	Val	Cys 55	Leu	Leu	Pro	Leu	Pro 60	Ser	Ala	Arg	Gly
	Asp 65	Pro	Glu	Trp	Ser	Gly 70	Gly	Phe	Phe	Phe						
40	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: "	710:							
45				(A) L B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	35 a no a lin	mino cid ear	aci		. 71	٥.			
50	Asp 1	Asp		_	Phe						•			Ala	Lys 15	His
50		Ile	Leu		5 Pro	Glu	Gly	Leu			Gly	Ala	Val			Ser
55	Ser	Lys	Lys 35		Lys	Arg	Asp	Leu 40	25 Ile	Asp	Asn	Ser	Phe 45		Arg	Tyr
60	Thr	Phe 50	Asn		Asp	Glu	Gly 55	Glu	Leu	Pro	Glu	Trp 60			Gln	Glui

```
Glu Lys Gln His Arg Ile Arg Gln Leu Pro Val Gly Lys Lys Glu Val
                       70
                                         75
     Glu His Tyr Arg Lys Arg Trp Arg Glu Ile Asn Ala Arg Pro Ile Xaa
 5
     100
                                 105
10
     Leu Glu Gln Thr Arg Lys Lys Ala Glu Ala Val Val Asn Thr Val Asp
                          120
     Ile Xaa Arg Thr Arg Glu Ser
15
     (2) INFORMATION FOR SEQ ID NO: 711:
20
            (i) SEQUENCE CHARACTERISTICS:
                  (A) LENGTH: 50 amino acids
                  (B) TYPE: amino acid
                  (D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 711:
25
     Asp Asp Asp Gly Phe Glu Ile Val Pro Ile Glu Asp Pro Ala Lys His
                                     10
     Arg Ile Leu Asp Pro Glu Gly Leu Ala Leu Gly Ala Val Ile Ala Ser
30
     Ser Lys Lys Ala Lys Arg Asp Leu Ile Asp Asn Ser Phe Asn Arg Tyr
                               40
35
     Thr Phe
40
     (2) INFORMATION FOR SEQ ID NO: 712:
            (i) SEQUENCE CHARACTERISTICS:
                  (A) LENGTH: 51 amino acids
                  (B) TYPE: amino acid
45
                  (D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 712:
     Lys Arg Trp Arg Glu Ile Asn Ala Arg Pro Ile Xaa Xaa Xaa Xaa Xaa
       1
                5
                                     10
50
     Arg Lys Lys Ala Glu Ala Val Val Asn Thr Val Asp Ile Xaa Arg Thr
55
                               40
     Arg Glu Ser
         50
```

708

5			(i) 5 (xi)	0	A) L B) T D) T	engi YPE : OPOL	H: 2 ami OGY:	16 ar no ac line	mino cid ear	aci		: 71 :	3:			
10	Met 1	Ile	Lys	Asp	Lys 5	Gly	Arg	Ala	Arg	Thr 10	Ala	Leu	Thr	Ser	Ser 15	Gln
15	Pro	Ala	His	Leu 20	Cys	Pro	Glu	Asn	Pro 25	Leu	Leu	His	Leu	Lys 30	Ala	Ala
13	Val	Lys	Glu 35	Lys	Lys	Arg	Asn	Lys 40	Lys	Lys	Lys	Thr	Ile 45	Gly	Ser	Pro
20	Lys	Arg 50	Ile	Gln	Ser	Pro	Leu 55	Asn	Asn	Lys	Leu	Leu 60	Asn	Ser	Pro	Ala
	Lys 65	Thr	Leu	Pro	Gly	Ala 70	Суз	Gly	Ser	Pro	Gln 75	Lys	Leu	Ile	Asp	Gl _y 80
25	Phe	Leu	Lys	His	Glu 85	Gly	Pro	Pro	Ala	Glu 90	Lys	Pro	Leu	Glu	Glu 95	Leu
30	Ser	Ala	Ser	Thr 100	Ser	Gly	Val	Pro	Gly 105	Leu	Ser	Ser	Leu	Gln 110	Ser	Ası
	Pro	Ala	Gly 115	Cys	Val	Arg	Pro	Pro 120	Ala	Pro	Asn	Leu	Ala 125	Gly	Ala	Va.
35		130				_	135					140				
	145		Pro			150					155					160
40			lle		165					170					175	
45			: Lys	180	1				185					190		
	Ala	Phe	195		Ile	: Leu	. Asp	Asn 200	Val	Gln	Val	Val	Leu 205		Gln	Th
50	Tyr	210 210	y Ser)	Thr	Leu	Lys	Val 215									
55	(2)	INE	FORMA	TION	FOR	SEC) ID	NO:	714:							
			(i)		(A) 1	LENG	rH: !	TERIS 52 an	nino		ds					
60			(xi)					: lir		SEO I	ED NO): 7 1	14:			

(2) INFORMATION FOR SEQ ID NO: 713:

WO 98/54963

PCT/US98/11422

•	Met Ile Lys Asp Lys Gly Arg Ala Arg Thr Ala Leu Thr Ser Ser Gln 1 5 10 15
5	Pro Ala His Leu Cys Pro Glu Asn Pro Leu Leu His Leu Lys Ala Ala 20 25 30
10	Val Lys Glu Lys Lys Arg Asn Lys Lys Lys Lys Thr Ile Gly Ser Pro 35 40 45
	Lys Arg Ile Gln 50
15	(2) INFORMATION FOR SEQ ID NO: 715:
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 100 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 715:
25	Lys Arg Ile Gln Ser Pro Leu Asn Asn Lys Leu Leu Asn Ser Pro Ala 1 5 10 15
	Lys Thr Leu Pro Gly Ala Cys Gly Ser Pro Gln Lys Leu Ile Asp Gly 20 25 30
30	Phe Leu Lys His Glu Gly Pro Pro Ala Glu Lys Pro Leu Glu Glu Leu 35 40 45
35	Ser Ala Ser Thr Ser Gly Val Pro Gly Leu Ser Ser Leu Gln Ser Asp 50 55 60
	Pro Ala Gly Cys Val Arg Pro Pro Ala Pro Asn Leu Ala Gly Ala Val 65 70 75 80
40	Glu Phe Asn Asp Val Lys Thr Leu Leu Arg Glu Trp Ile Thr Thr Ile 85 90 95
	Ser Asp Pro Met 100
45	(2) INFORMATION FOR SEQ ID NO: 716:
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 74 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 716:
55	Thr Ile Ser Asp Pro Met Glu Glu Asp Ile Leu Gln Val Val Lys Tyr 1 5 10 15
60	Cys Thr Asp Leu Ile Glu Glu Lys Asp Leu Glu Lys Leu Asp Leu Val 20 25 30

WO 98/54963

PCT/US98/11422

710

Ile Lys Tyr Met Lys Arg Leu Met Gln Gln Ser Val Glu Ser Val Trp 35 Asn Met Ala Phe Asp Phe Ile Leu Asp Asn Val Gln Val Val Leu Gln 5 55 Gln Thr Tyr Gly Ser Thr Leu Lys Val Thr 70 10 (2) INFORMATION FOR SEQ ID NO: 717: (i) SEQUENCE CHARACTERISTICS: 15 (A) LENGTH: 18 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 717: 20 Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Asn Cys 5 10 Glu Pro 25 (2) INFORMATION FOR SEQ ID NO: 718: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 718: 35 Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Asn Cys 5 10 Glu Pro 40 (2) INFORMATION FOR SEQ ID NO: 719: 45 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 719: Pro Gln Pro Ser Asn Phe Pro Thr Thr Val Arg Asn Leu Pro Tyr Ser 5 10 55 Gly Ala Gly Ala Gln Pro Pro Pro Ser Asn Cys 20 60 (2) INFORMATION FOR SEQ ID NO: 720:

c			(i) :	C	A) L B) T	ENGT YPE:	H: 1 ami	34 au no a	mino cid		ds					
5			(xi)	SEQ		OPOLA E DES				EQ II	ои с	: 720	0:			
10	Met 1	Ala	Ser	Ser	Val 5	Pro	Ala	Gly	Gly	His 10	Thr	Arg	Ala	Gly	Gly 15	Ile
10	Phe	Leu	Ile	Gly 20	Lys	Leu	Asp	Leu	Glu 25	Ala	Ser	Leu	Phe	Lys 30	Ser	Phe
15	Gln	Trp	Leu 35	Pro	Phe	Val	Leu	Arg 40	Lys	ГЛЗ	Cys	Asn	Phe 45	Phe	Cys	Trp
	Asp	Ser 50		Ala	His	Ser	Leu 55	Pro	Leu	His	Pro	Leu 60	Ser	Ala	Ser	Cys
20	Ser 65	Ala	Pro	Ala	Cys	His 70	Ala	Ser	Asp	Thr	His 75	Leu	Leu	Tyr	Pro	Ser 80
25	Thr	Arg	Ala	Leu	Cys 85	Pro	Ser	Ile	Phe	Ala 90	Trp	Leu	Val	Ala	Pro 95	His
23	Ser	Val	Phe	Arg 100	Thr	Asn	Ala	Pro	Gly 105	Pro	Thr	Pro	Ser	Ser 110	Gln	Ser
30	Ser	Pro	Val 115	Phe	Pro	Val	Phe	Pro 120	Val	Ser	Phe	Met	Ala 125	Leu	Ile	Val
	Суѕ	Хаа 130		Val	Cys	Cys								٠.		
35																
	(2)	INF		TION			•									
40			(i)		(A) I (B) 7	ENGI	H: 7	1 an	nino cid		is					
			(xi)	SEC		OPOI E DE				EQ I	D NO): 7 2	1:			
45	Met 1		. Ser	Ser	Val		Ala	Gly	Gly	His		Arg	Ala	Gly	Gly 15	
50	Phe	Leu	ı Ile	Gly 20		Leu	Asp	Leu	Glu 25		Ser	Leu	Phe	Lys 30		Phe
50	Glm	Tr	Let 35	ı Pro	Phe	. Val	Leu	Arg		Lys	Cys	: Asn	Phe 45		Cys	Trp
55	Asp	Se: 5(Ala	His	Ser	Leu 55		Leu	His	Pro	Leu 60		Ala	Ser	Cys
	Ser 65		a Pro	Ala	Cys	His 70		L								

	(2) 1	NFC	RMAT	ION	FOR	SEQ	ID N	0: 7	22:							
5			(i) S (xi)	() (I	A) LI 3) Ti 0) T(ENGTI PE: OPOLO	i: 40 amir OGY:	o ami no ac line	ino a cid ear	acid		722	2:			
10	Phe I	Ala	Trp	Leu	Val 5	Ala	Pro	His	Ser	Val 10	Phe	Arg	Thr	Asn	Ala 15	Pro
15	Gly I			20					25					30	Phe	Pro
20	vui .	JC1	35			200		40	-,-				45	-3-		
20	(2)		ORMAI													
25			(xi)	(A) L B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	34 a no a lin	mino cid ear	aci		: 72	3:			
30	Met .	Ala	Ser	Ser	Val 5	Pro	Ala	Gly	Gly	His 10	Thr	Arg	Ala	Gly	Gly 15	Ile
	Phe	Leu	Ile	Gly 20	Lys	Leu	Asp	Leu	Glu 25	Ala	Ser	Leu	Phe	Lys 30	Ser	Phe
35	Gln	Trp	Leu 35	Pro	Phe	Val	Leu	Arg 40	Lys	Lys	Cys	Asn	Phe 45	Phe	Cys	Trp
40	Asp	Ser 50		Ala	His	Ser	Leu 55	Pro	Leu	His	Pro	Leu 60	Ser	Ala	Ser	Cys
	Ser 65	Ala	Pro	Ala	Cys	His 70		Ser	Asp	Thr	His 75	Leu	Leu	Tyr	Pro	Ser 80
45	Thr	Arg	Ala	Leu	Cys 85	Pro	Ser	Ile	Phe	Ala 90		Leu	Val	Ala	Pro 95	His
	Ser	Val	Phe	Arg 100		Asn	Ala	Pro	Gly 105		Thr	Pro	Ser	Ser 110		Ser
50	Ser	Pro	Val 115		Pro	Val	Phe	Pro 120	Val	Ser	Phe	Met	Ala 125		Ile	Val
55	Суз	Xaa 130	Leu	Val	Cys	Cys										
	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	724:							

(i) SEQUENCE CHARACTERISTICS:

•	(A) LENGTH: 286 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 724:															
_		1	(xi)	SEQU	JENCI	E DES	SCRIE	MOITS	i: SE	O II) NO:	724	1:			
5	Met 1	Ala	Met	Glu	Gly 5	Tyr	Trp	Arg	Phe	Leu 10	Ala	Leu	Leu	Gly	Ser 15	Ala
10	Leu	Leu	Val	Gly 20	Phe	Leu	Ser	Val	Ile 25	Phe	Ala	Leu	Val	Trp 30	Val	Leu
	His	Tyr	Arg 35	Glu	Gly	Leu	Gly	Trp 40	Asp	Gly	Ser	Ala	Leu 45	Glu	Phe	Asn
15	Trp	His 50	Pro	Val	Leu	Met	Val 55	Thr	Gly	Phe	Val	Phe 60	Ile	Gln	Gly	Ile
20	Ala 65	Ile	Ile	Val	Туг	Arg 70	Leu	Pro	Trp	Thr	Trp 75	Lys	Cys	Ser	Lys	Leu 80
20	Leu	Met	Lys	Ser	Ile 85	His	Ala	Gly	Leu	Asn 90	Ala	Val	Ala	Ala	Ile 95	Leu
25	Ala	Ile	Ile	Ser 100	Val	Val	Ala	Val	Phe 105	Glu	Asn	His	Asn	Val 110	Asn	Asn
	Ile	Ala	Asn 115	Met	Tyr	Ser	Leu	His 120	Ser	Trp	Val	Gly	Leu 125	Ile	Ala	Val
30	Ile	Cys 130		Leu	Leu	Gln	Leu 135	Leu	Ser	Gly	Phe	Ser 140	Val	Phe	Leu	Leu
35	Pro 145	Trp	Ala	Pro	Leu	Ser 150		Arg	Ala	Phe	Leu 155	Met	Pro	Ile	His	Val 160
	Tyr	Ser	Gly	Ile	Val 165		Phe	Gly	Thr	Val 170		Ala	Thr	Ala	Leu 175	Met
40	Gly	Leu	Thr	Glu 180	-	Leu	Ile	Phe	Ser 185		Arg	Asp	Pro	Ala 190		Ser
٠.	Thr	Phe	Pro 195		Glu	Gly	Val	Phe 200		Asn	Thr	Leu	Gly 205		Leu	Ile
45	Leu	Val 210		Gly	Ala		1le 215		Trp	Ile		Thr 220		Pro	Gln	Tr
50	Lys 225	Arg	Pro	Lys	Glu	230		Ser	Thr	Ile	Leu 235		Pro	Asn	Gly	G1 ₃
	Thr	Glu	Gln	Gly	Ala 245		Gly	Ser	Met	250		Тут	Ser	Gly	255	
55	Met	: Asp	Lys	Ser 260		Ser	Glu	Leu	Asn 265		Glu	V al	. Ala	Ala 270		Ly:
	Arg	J Asr	275		Leu	ı Ası	Glu	Ala 280	_	Gln	Arg	Ser	Thr 285		:	

714

5				(ENCE A) L B) T D) T UENC	ENGT YPE: OPOL	H: 4 ami OGY:	3 am no a lin	ino cid ear	acid		: 72	5 :			
10	Pro 1	Gly	Arg	Ala	Gly 5	Pro	Ser	Pro	Gly	Leu 10	Ser	Leu	Gln	Leu	Pro 15	Ala
15	Glu	Pro	Gly	His 20	Pro	Ala	Gly	Asn	Leu 25	Ala	Pro	Leu	Thr	Ser 30	Arg	Pro
	Gln	Pro	Leu 35	Cys	Arg	Ile	Pro	Ala 40	Val	Pro	Gly					
20	(2)	INF	ORMAT	rion	FOR	SEQ	ID I	NO: 1	726:							
25			(i) : (xi)	(A) L B) T D) T	ENGT YPE : OPOL	H: 4 ami OGY:	24 a no a lin	mino cid ear	aci		: 72	6 :			
30	Met 1	Lys	Leu	Leu	Gly 5	Glu	Суз	Ser	Ser	Ser 10	Ile	Asp	Ser	Val	Lys 15	Arg
	Leu	Glu	His	Lys 20	Leu	Lys	Glu	Glu	Glu 25	Glu	Ser	Leu	Pro	Gly 30	Phe	Val
35	Asn	Ļeu	His 35	Ser	Thr	Glu	Thr	Gln 40	Thr	Ala	Gly	Val	Ile 45	Asp	Arg	Trp
40	Glu	Leu 50	Leu	Gln	Ala	Gln	Ala 55	Leu	Ser	Lys	Glu	Leu 60	Arg	Met	Lys	Gln
٠.	Asn 65	Leu	Gln	Lys	Trp	Gln 70	Gln	Phe	Asn	Ser	Asp 75	Leu	Asn	Ser	Ile	Trp 80
45	Ala	Trp	Leu	Gly	Asp 85	Thr	Glu	Glu	Glu	Leu 90	Glu	Gln	Leu	Gln	Arg 95	Leu
	Glu	Leu	Ser	Thr 100	Asp	Ile	Gln	Thr	Ile 105	Glu	Leu	Gln	Ile	Lys 110	Lys	Leu
50	Lys	Glu	Leu 115	Gln	Lys	Ala	Val	Asp 120	His	Arg	Lys	Ala	Ile 125	Ile	Leu	Ser
55	Ile	Asn 130	Leu	Суѕ	Ser	Pro	Glu 135	Phe	Thr	Gln	Ala	Asp 140	Ser	Lys	Glu	Ser
	Arg 145	Asp	Leu	Gln	Asp	Arg 150	Leu	Xaa	Gln	Met	Asn 155	Gly	Arg	Trp	Asp	Arg 160
60	Val	Суз	Ser	Leu	Leu 165	Glu	Glu	Trp	Arg	Gly 170	Leu	Leu	Gln	Asp	Ala 175	Leu

(2) INFORMATION FOR SEQ ID NO: 725:

	Met	Gln	Cys	Gln 180	Gly	Phe	His	Glu	Met 185	Ser	His	Gly	Leu	Leu 190	Leu	Met
5	Leu	Glu	Asn 195	Ile	Asp	Arg	Arg	Lys 200	Asn	Glu	Ile	Val	Pro 205	Ile	Asp	Ser
10	Asn	Leu 210	Asp	Ala	Glu	Ile	Leu 215	Gln	Asp	His	His	Lys 220	Gln	Leu	Met	Gln
10	Ile 225	Lys	His	Glu	Leu	Leu 230	Glu	Ser	Gln	Leu	Arg 235	Val	Ala	Ser	Leu	Gln 240
15	Asp	Met	Ser	Cys	Gln 245	Leu	Leu	Val	Asn	Ala 250	Glu	Gly	Thr	Asp	Cys 255	Leu
	Glu	Ala	Lys	Glu 260	Lys	Val	His	Val	Ile 265	Gly	Asn	Arg	Leu	Lys 270	Leu	Leu
20	Leu	Lys	Glu 275	Val	Ser	Arg	His	Ile 280	Lys	Glu	Leu	Glu	Lys 285	Leu	Leu	Asp
25	Val	Ser 290	Ser	Ser	Gln	Gln	Asp 295	Leu	Ser	Ser	Trp	Ser 300	Ser	Ala	Asp	Glu
	Leu 305	Asp	Thr	Ser	Gly	Ser 310	Val	Ser	Pro	Xaa	Ser 315	Gly	Arg	Ser	Thr	Pro 320
30	Asn	Arg	Gln	Lys	Thr 325	Pro	Arg	Gly	Lys	Cys 330	Ser	Leu	Ser	Gln	Pro 335	Gly
	Pro	Ser	Val	Ser 340	Ser	Pro	His	Ser	Arg 345	Ser	Thr	Lys	Gly	Gly 350	Ser	Asp
35	Ser	Ser	Leu 355	Ser	Glu	Pro	Xaa	Pro 360	Gly	Arg	Ser	Gly	Arg 365	Gly	Phe	Leu
40	Phe	Arg 370		Leu	Arg	Ala	Ala 375	Leu	Pro	Leu	Gln	Leu 380	Leu	Leu	Leu	Leu
,	Leu 385		Gly	Leu	Ala	Cys 390		Val	Pro	Met	Ser 395	Glu	Glu	Asp	Tyr	Ser 400
45	Cys	Ala	Leu	Ser	Asn 405	Asn	Phe	Ala	Arg	Ser 410	Phe	His	Pro	Met	Leu 415	Arg
50	Tyr	Thr	Asn	Gly 420	Pro	Pro	Pro	Leu								
50	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	727 :							
55				(ENCE (A) I (B) I (D) I	PPE:	TH: 1 ami	ino a lno a	mino cid ear	aci): 72	: 7 :			
60	Mo-			I or	Cly	c1	0	°	Co~	· Co-	71~	. x	· Ca-	1751	7	7~~

(2) INFORMATION FOR SEQ ID NO: 729:

	1				5					10					15	
5	Leu	Glu	His	Lys 20	Leu	Lys	Glu	Glu	Glu 25	Glu	Ser	Leu	Pro	Gly 30	Phe	Val
5	Asn	Leu	His 35	Ser	Thr	Glu	Thr	Gln 40	Thr	Ala	Gly	Val	Ile 45	Asp	Arg	Trp
10	Glu	Leu 50	Leu	Gln	Ala	Gln	Ala 55	Leu	Ser	Lys	Glu	Leu 60	Arg	Met	Lys	Gln
	Asn 65	Leu	Gln	Lys	Trp	Gln 70	Gln	Phe	Asn	Ser	Asp 75	Leu	Asn	Ser	Ile	Trp 80
15	Ala	Trp	Leu	Gly	Asp 85	Thr	Glu	Glu	Glu	Leu 90	Glu	Gln	Leu	Gln	Arg 95	Leu
20	Glu	Leu	Ser	Thr 100	Asp	Ile	Gln	Thr	Ile 105	Glu	Leu	Gln	Ile	Lys 110		
	(2)	INFO	ORMA	rion	FOR	SEQ	ID 1	10:	728:							
25			(i) :	(A) L B) T	ENGI YPE :		36 a no a	mino cid	: aci	ds					
30			(xi)		-					EQ I	D NO	: 72	8:			
	Lys 1	Leu	Lys	Glu	Leu 5	Gln	Lys	Ala	Val	Asp 10	His	Arg	Lys	Ala	Ile 15	Ile
35	Leu	Ser	Ile	Asn 20	Leu	Cys	Ser	Pro	Glu 25		Thr	Gln	Ala	Asp 30	Ser	Lys
	Glu	Ser	Arg 35	Asp	Leu	Gln	Asp	Arg 40		Xaa	Gln	Met	Asn 45		Arg	Trp
40	Asp	Arg 50		Суз	Ser	Leu	Leu 55		Glu	Trp	Arg	Gly 60	Leu	Leu	Gln	Asp
45	65					70					75					Leu 80
	Leu	Met	Leu	Glu	Asn 85		Asp	Arg	Arg	Lys 90		Glu	Ile	Val	Pro 95	Ile
50	Asp	Ser	Asn	Leu 100		Ala	Glu	Ile	Leu 105		Asp	His	His	Lys 110		Leu
	Met	Gln	11s		His	Glu	Leu	120	_	Ser	Gln	Leu	Arg 125		Ala	Ser
55	Leu	Gln 130	Asp	Met	Ser	Cys	Gln 135		ı							

	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 105 amino acids (B) TYPE: amino acid															
5			(xi)		D) T	OPOL	OGY:	line	ear	D II	OM C	: 729):			
10	Gln 1	Ąsp	Met	Ser	Cys 5	Gln	Leu	Leu	Val	Asn 10	Ala	Glu	Gly	Thr	Asp 15	Суз
10	Leu	Glu	Ala	Lys 20	Glu	Lys	Val	His	Val 25	Ile	Gly	Asn	Arg	Leu 30	Lys	Leu
15	Leu	Leu	Lys 35	Glu	Val	Ser	Arg	His 40	Ile	Lys	Glu	Leu	Glu 45	Lys	Leu	Leu
	Asp	Val 50	Ser	Ser	Ser	Gln	Gln 55	Asp	Leu	Ser	Ser	Trp 60	Ser	Ser	Ala	Asp
20	Glu 65	Leu	Asp	Thr	Ser	Gly 70	Ser	Val	Ser	Pro	Xaa 75	Ser	Gly	Arg	Ser	Thr 80
25	Pro	Asn	Arg	Gln	Lys 85	Thr	Pro	Arg	Gly	Lys 90	Cys	Ser	Leu	Ser	Gln 95	Pro
	Gly	Pro	Ser	Val 100	Ser	Ser	Pro	His	Ser 105							
30	(2)	INF	ORMA'	rion	FOR	SEO	ID I	NO: 1	730:							
	(2) INFORMATION FOR SEQ ID NO: 730: (i) SEQUENCE CHARACTERISTICS:															
35 -				(B) T	ENGT YPE: OPOL	ami	no a	cid	acid	s					
			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 73	0:			
40	Asp 1	Ser	Ser	Leu	Ser 5	Glu	Pro	Xaa	Pro	Gly 10	Arg	Ser	Gly	Arg	Gly 15	Phe
	Leu	Phe	Arg	Val 20	Leu	Arg	Ala	Ala	Leu 25	Pro	Leu	Gln	Leu	Leu 30	Leu	Leu
45	Leu	Leu	Ile 35	Gly	Leu	Ala	Cys	Leu 40	Val	Pro	Met	Ser	Glu 45	Glu	Asp	Tyr
50	Ser	Cys 50		Leu	Ser	Asn	Asn 55	Phe	Ala	Arg	Ser	Phe 60	His	Pro	Met	Leu
	Arg 65	_	Thr	Asn	Gly	Pro 70	Pro	Pro	Leu							
55	(2)	INF	ORMA	TION	FOR	SEQ	ID :	NO: '	731:							
			(i)	SEQU							_					
60				-		ENGI YPE:				acid	ls					

(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 731:													
5	Met Lys Leu Leu Ile Cys Gly Asn Tyr Leu Ala Pro Ser His 1 5 10	s Ser Glu 15											
	Ser Ser Arg Arg Cys Cys Leu Leu Cys Phe Tyr Pro Leu Cys 20 25 30												
10	Ile Asn Phe Gly Met Lys Val Phe Leu Ser Met Pro Phe Leu 35 40 45	u Val Leu											
15	Phe Gln Ser Leu Ile Gln Glu Asp 50 55												
20	(2) INFORMATION FOR SEQ ID NO: 732: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 271 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear												
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 732: Arg Ile Leu Leu Val Lys Tyr Ser Ala Asn Glu Glu Asn Ly 1 5 10	rs Tyr Asp 15											
30		30											
	Phe Cys Val Leu Val Ser Phe Cys Val Ile Lys Lys Asp Hi 35 40 45	s Gln Ser											
35	Arg Asn Leu Lys Tyr Ala Ser Trp Lys Glu Phe Ser Asp Ph 50 55 60	e Met Lys											
40	Trp Ser Ile Pro Ala Phe Leu Tyr Phe Leu Asp Asn Leu II 65 70 75	80											
	Tyr Val Leu Ser Tyr Leu Gln Pro Ala Met Ala Val Ile Pi 85 90	ne Ser Asn 95											
45	Phe Ser Ile Ile Thr Thr Ala Leu Leu Phe Arg Ile Val Leu 100 105 1:	eu Lys Xaa 10											
	Arg Leu Asn Trp Ile Gln Trp Ala Ser Leu Leu Thr Leu Pl 115 120 125	ne Leu Ser											
50	Ile Val Ala Leu Thr Ala Gly Thr Lys Thr Leu Gln His A 130 135 140	sn Leu Ala											
55	Gly Arg Gly Phe His His Asp Ala Phe Phe Ser Pro Ser A 145 150 155	160											
	Leu Leu Phe Arg Asn Glu Cys Pro Arg Lys Asp Asn Cys T 165 170	hr Ala Lys 175											
60	Glu Trp Thr Phe Pro Glu Ala Lys Trp Asn Thr Thr Ala A 180 185 1	rg Val Phe 90											

	Ser	His	Ile 195	Arg	Leu	Gly	Met	Gly 200	His	Val	Leu	Ile	Ile 205	Val	Gln	Cys
5	Phe	Ile 210	Ser	Ser	Met	Ala	Asn 215	Ile	Tyr	Asn	Glu	Lys 220	Ile	Leu	Lys	Glu
10	Gly 225	Asn	Gln	Leu	Thr	Glu 230	Xaa	Ile	Phe	Ile	Gln 235	Asn	Ser	Lys	Leu	Туг 240
	Phe	Phe	Gly	Ile	Leu 245	Phe	Asn	Gly	Leu	Thr 250	Leu	Gly	Leu	Gln	Arg 255	Ser
15	Asn	Arg	Asp	Gln 260	Ile	Lys	Asn	Суз	Gly 265	Phe	Phe	Tyr	Gly	His 270	Ser	
20	(2)	INF				CHA:	RACT H: 9	ERIS	TICS ino		s					
25			(xi)		D) T	OPOL	OGY:	lin	ear	EQ I	D NO	: 73	3:			
	Asn 1	Ser	Val	Pro	Asn 5	Leu	Gln	Thr	Leu	Ala 10	Val	Leu	Thr	Glu	Ala 15	Ile
30	Gly	Pro	Glu	Pro 20	Ala	Ile	Pro	Arg	Хаа 25	Pro	Arg	Glu	Pro	Pro 30	Val	Ala
35	Thr	Ser	Thr 35	Pro	Ala	Thr	Pro	Ser 40	Ala	Gly	Pro	Gln	Pro 45	Leu	Pro	Thr
	Gly	Thr 50		Leu	Val	Pro	Gly 55	Gly	Pro	Ala	Pro	Pro 60	Cys	Leu	Gly	Glu
40	Ala 65		Ala	Leu	Leu	Leu 70	Pro	Pro	Cys	Arg	Pro 75	Ser	Leu	Thr	Ser	Cys 80
	Phe	Trp	Ser	Pro	Arg 85	Pro	Ser	Pro	Trp	Lys 90	Glu	Thr	Gly	Val		
45	(2)	INF	ORMA	TION	FOR	SEO	ID i	NO: '	734:							
50			(i)	SEQU (ENCE (A) I (B) I (D) I	CHA ENGI YPE:	RACT H: 4 ami OGY:	ERIS 0 am no a lin	TICS ino cid ear	acid		: 73	4:			
55	Ala 1		Gln	Leu	Ala 5	Phe	Tyr	Pro	Asp	Ala 10	Val	Glu	Glu	Trp	Leu 15	Glu
60	Glu	Asn	Val	His 20	Pro	Ser	Leu	Gln	Arg 25	Leu	Gln	Хаа	Leu	Leu 30	Gln	Asp

720

Leu Ser Glu Val Ser Ala Pro Pro 35 5 (2) INFORMATION FOR SEQ ID NO: 735: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids 10 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 735: Cys His Pro Pro Ala Leu Ala Gly Thr Leu Leu Arg Thr Pro Glu Gly 15 Arg Ala His Ala Arg Gly Leu Leu Glu Ala Gly Gly Ala 20 20 (2) INFORMATION FOR SEQ ID NO: 736: (i) SEQUENCE CHARACTERISTICS: 25 (A) LENGTH: 59 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 736: 30 Gly Ser Ser Ser Thr Arg Ser Trp Phe Ser Thr Ser Ser Pro Gln Arg 10 Ser Ala Ser Trp His Ser Gly Ala Pro Ser Cys Arg Ser Trp Arg Leu 20 25 30 35 Pro Cys Ser Trp Leu Ser Thr Arg Met Pro Trp Arg Ser Gly Trp Arg Lys Thr Cys Thr Pro Ala Cys Ser Gly Cys Lys 40 (2) INFORMATION FOR SEQ ID NO: 737: 45 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 247 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 737: Met Arg Pro Asp Trp Lys Ala Gly Ala Gly Pro Gly Gly Pro Pro Gln 5 55 Lys Pro Ala Pro Ser Ser Gln Arg Lys Pro Pro Ala Arg Pro Ser Ala Ala Ala Ala Ile Ala Val Ala Ala Ala Glu Glu Glu Arg Arg Leu 40 60

•	Arg	Gln 50	Arg	Asn	Arg	Leu	Arg 55	Leu	Glu	Glu	Asp	Lys 60	Pro	Ala	Val	Glu
5	Arg 65	Cys	Leu	Glu	Glu	Leu 70	Val	Phe	Gly	Asp	Val 75	Glu	Asn	Asp	Glu	Asp 80
	Ala	Leu	Leu	Arg	Arg 85	Leu	Arg	Gly	Pro	Arg 90	Val	Gln	Glu	His	Glu 95	Asp
10	Ser	Gly	Asp	Ser 100	Glu	Val	Glu	Asn	Glu 105	Ala	Lys	Gly	Asn	Phe 110	Pro	Pro
15	Gln	Lys	Lys 115	Pro	Val	Trp	Val	Asp 120	Glu	Glu	Asp	Glu	Asp 125	Glu	Glu	Met
	Val	Asp 130	Met	Met	Asn	Asn	Arg 135	Phe	Arg	Lys	Asp	Met 140	Met	Lys	Asn	Ala
20	Ser 145	Glu	Ser	Lys	Leu	Ser 150	Lys	Asp	Asn	Leu	Lys 155	Lys	Arg	Leu	Lys	Glu 160
	Glu	Phe	Gln	His	Ala 165	Met	Gly	Gly	Val	Pro 170	Ala	Trp	Ala	Glu	Thr 175	Thr
25	Lys	Arg	Lys	Thr 180	Ser	Ser	Asp	Asp	Glu 185	Ser	Glu	Glu	Asp	Glu 190	Asp	Asp
30	Leu	Leu	Gln 195	Arg	Thr	Gly	Asn	Phe 200	Ile	Ser	Thr	Ser	Thr 205	Ser	Leu	Pro
	Arg	Gly 210	Ile	Leu	Lys	Met	Lys 215	Asn	Cys	Gln	His	Ala 220	Asn	Ala	Glu	Arg
35	Pro 225	Thr	Val	Ala	Arg	Ile 230	Ser	Ile	Cys	Ala	Val 235	Pro	Ser	Arg	Cys	Thr 240
	Asp	Cys	Asp	Gly	Cys 245	Trp	Asp									
40																
	(2)		ORMAT													
45				() () ()	A) Li B) T D) T	ENGTI YPE : OPOLA	H: 1: ami: OGY:	80 ar no ac line	mino cid ear	acio						
50			(xi)													
50	Cys 1	Leu	Glu	Glu	Leu 5	Val	Phe	Gly	Asp	Val 10	Glu	Asn	Asp	Glu	Asp 15	Ala
55	Leu	Leu	Arg	Arg 20	Leu	Arg	Gly	Pro	Arg 25	Val	Gln	Glu	His	Glu 30	Asp	Ser
	Gly	Asp	Ser 35	Glu	Val	Glu	Asn	Glu 40	Ala	Lys	Gly	Asn	Phe 45	Pro	Pro	Gln
60	Lys	Lys 50	Pro	Val	Trp	Val	Asp 55	Glu	Glu	Asp	Glu	Asp 60	Glu	Glu	Met '	Val

	AS)	o me	c Me	C ASI	n Ası	1 Arg		e Arg	J Lys	s Asp	75		. Lys	Asn	Ala	Ser 80
5	Gl	ı Se:	r Ly:	s Lev	Ser 85	Lys	s Ası	Asr	Leu	Lys 90		Arg	Leu	Lys	Glu 95	
10	Pho	e Glı	n His	100	Met	: Gly	/ Gly	/ Val	. Pro	Ala	Trp	Ala	Glu	Thr 110		Lys
	Arg	J Ly:	5 Thi 119	Ser	Ser	Asp) Asp	Glu 120		Glu	Glu	Asp	Glu 125		Asp	Leu
15	Let	130	ı Arg	Thr	Gly	Asn	Phe 135		Ser	Thr	Ser	Thr 140		Leu	Pro	Arg
	Gl ₃ 145	/ Ile	e Leu	Lys	Met	Lys 150		Cys	Gln	His	Ala 155	Asn	Ala	Glu	Arg	Pro 160
20	The	Va]	l Ala	Arg	11e 165	Ser	Ile	Cys	Ala	Val 170	Pro	Ser	Arg	Суз	Thr 175	Asp
25	Cys	Asp	Gly	Cys 180									-			
	(2)	INF	ORMA	TION	FOR	SEO	ID:	NO:	739:							
30				SEQU	ENCE	СНА	RACT	ERIS	TICS	: aci	ds					
			(xi)	((B) 1 (D) 1	YPE: OPOI	ami OGY:	no a	cid	EQ II		. 73	g -			
35	Leu 1	Lys								Glu 10				Asp	Gly 15	Ser
40	Phe	Leu	Leu	Ile 20	Asn	Gly	Ile	Ala	Gly 25	Tyr	Leu	His	Leu	Leu 30	Ala	Met
٠.	Lys	Thr	Lys 35	Glu	Leu	Ile	Gly	Ser 40	Met	Lys	Ile	Asn	Gly 45	Arg	Val	Ala
45	Ala	Ser 50	Thr	Phe	Ser	Ser	Asp 55	Ser	Lys	Lys	Val	Tyr 60	Ala	Ser	Ser	Gly
50	Asp 65	Gly	Glu	Val	Tyr	Val 70	Trp	Asp	Val	Asn	Ser 75	Arg	Lys	Cys	Leu	Asn 80
	Arg	Phe	Val	Asp	Glu 85	Gly	Ser	Leu	Tyr	Gly 90	Leu	Ser	Ile		Thr 95	Ser
55	Arg	Asn	Gly	Gln 100	Tyr	Val	Ala	Суз	Gly 105	Ser	Asn	Суз	Gly	Val 110	Val	Asn
	Ile	Тут	Asn 115	Gln	Asp	Ser	Cys	Leu 120	Gln	Glu	Thr		Pro 125	Lys	Pro	Ile
60	Lys	Ala	Ile	Met	Asn	Leu	Val	Thr	Glv	Val	Thr	Ser	Leu	Thr	Phe	Asn

•	130	13	5	140	•
5	Pro Thr Thr G	lu Ile Leu Al 150	a Ile Ala Ser	Glu Lys Met 155	Lys Glu Ala 160
	Val Arg Leu V	al His Leu Pr 165	o Ser Cys Thr	Val Phe Ser	Asn Phe Pro
10	Val Ile Lys A 1	sn Lys Asn Il 80	e Ser His Val 1 185		Asp Phe Ser 190
	195		a Leu Gly Asn (200	Glu Lys Gly 1 205	Lys Ala Leu
15	Met Tyr Arg L 210	eu His His Tyr 219			
20	(2) INFORMATIO	ON FOR SEQ ID	NO: 740:		
25		(B) TYPE: am (D) TOPOLOGY	167 amino acida ino acid		
30	Lys Ile Asn Gl	y Arg Val Ala 5	A Ala Ser Thr P	he Ser Ser A	Asp Ser Lys 15
	Lys Val Tyr Al	a Ser Ser Gly O	/ Asp Gly Glu V 25	al Tyr Val T	Crp Asp Val 30
35	Asn Ser Arg Ly 35	s Cys Leu Asn	Arg Phe Val A 40	sp Glu Gly S 45	er Leu Tyr
40	50	55		60	
40	65	70		75	80
45	Glu Thr Asn Pr	85	90		95
	Val Thr Ser Le	u Thr Phe Asn O	Pro Thr Thr G		la Ile Ala 10
50	Ser Glu Lys Me 115	t Lys Glu Ala	Val Arg Leu Va 120	al His Leu P 125	ro Ser Cys
	Thr Val Phe Se 130	r Asn Phe Pro 135	Val Ile Lys As	sn Lys Asn I 140	le Ser His
55	Val His Thr Me 145	Asp Phe Ser 150	Pro Arg Ser G		la Leu Gly 160
60	Asn Glu Lys Gl	/ Lys Ala Leu 165			

	(2) INFORMATION FOR SEQ ID NO: 741: (i) SEQUENCE CHARACTERISTICS:															
5			(i) :	(A) L B) T	CHAI ENGT YPE: OPOL	H: 2 ami	46 a no a	mino cid		ds					
10			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D 190	: 74	1:			
	Met 1	Arg	Ile	Leu	Gln 5	Leu	Ile	Leu	Leu	Ala 10	Leu	Ala	Thr	Gly	Leu 15	Val
15	Gly	Gly	Glu	Thr 20	Arg	Ile	Ile	Lys	Gly 25	Phe	Glu	Суѕ	Lys	Leu 30	His	Ser
	Gln	Pro	Trp 35	Gln	Ala	Ala	Leu	Phe 40	Glu	Lys	Thr	Arg	Leu 45	Leu	Cys	Gly
20	Ala	Thr 50	Leu	Ile	Ala	Pro	Arg 55	Trp	Leu	Leu	Thr	Ala 60	Ala	His	Суз	Leu
25	Lys 65	Pro	Arg	Tyr	Ile	Val 70	His	Leu	Gly	Gln	His 75	Asn	Leu	Gln	Lys	Glu 80
	Glu	Gly	Cys	Glu	Gln 85	Thr	Arg	Thr	Ala	Thr 90	Glu	Ser	Phe	Pro	His 95	Pro
30	Gly	Phe	Asn	Asn 100	Ser	Leu	Pro	Asn	Lys 105	Asp	His	Arg	Asn	Asp 110	Ile	Met
	Leu	Val	Lys 115	Met	Ala	Ser	Pro	Val 120	Ser	Ile	Thr	Trp	Ala 125	Val	Arg	Pro
35	Leu	Thr 130	Leu	Ser	Ser	Arg	Cys 135	Val	Thr	Ala	Gly	Thr 140	Ser	Cys	Ser	Phe
40	Pro 145	Ala	Gly	Ala	Ala	Arg 150	Pro	Asp	Pro	Ser	Tyr 155	Ala	Cys	Leu	Thr	Pro 160
	Cys	Asp	Ala	Pro	Thr 165	Ser	Pro	Ser	Leu	Ser 170	Thr	Arg	Ser	Val	Arg 175	Thr
45	Pro	Thr	Pro	Ala 180	Thr	Ser	Gln	Thr	Pro 185	Trp	Cys	Val	Pro	Ala 190	Cys	Arg
	Lys	Gly	Ala 195	Arg	Thr	Pro	Ala	Arg 200	Val	Thr	Pro		Ala 205	Leu	Trp	Ser
50	Val	Thr 210	Ser	Leu	Phe	Lys	Ala 215	Leu	Ser	Pro	Gly	Ala 220	Arg	Ile	Arg	Val
55	Arg 225	Ser	Pro	Glu	Ser	Leu 230	Val	Ser	Thr	Arg	Lys 235	Ser	Ala	Asn	Met	Trp 240
=	Thr	Gly	Ser	Arg	Arg 245	Arg										

725

•	(2)	INF	ORMA	TION	FOR	SEQ	ID.	NO:	742:							
5				((A) I (B) I (D) I	ENGI YPE : YPOI	TH: 2 ami OGY:	28 a no a lir	mind cid ear	: o aci EQ I): 74	2 :			
10	Glu 1	Thr	Arg	Ile	Ile 5		Gly	Phe	Glu	Cys 10		Leu	His	Ser	Gln 15	
	Trp	Gln	Ala	Ala 20	Leu	Phe	Glu	Lys	Thr 25		Leu	Leu	Cys	Gly 30		Thi
15	Leu	Ile	Ala 35	Pro	Arg	Trp	Leu	Leu 40	Thr	Ala	Ala	His	Cys 45	Leu	Lys	Pro
20	Arg	Tyr 50	Ile	Val	His	Leu	Gly 55	Gln	His	Asn	Leu	Gln 60	Lys	Glu	Glu	Gly
20	Cys 65	Glu	Gln	Thr	Arg	Thr 70	Ala	Thr	Glu	Ser	Phe 75	Pro	His	Pro	Gly	Phe 80
25	Asn	Asn	Ser	Leu	Pro 85	Asn	Lys	Asp	His	Arg 90	Asn	Asp	Ile	Met	Leu 95	Val
	Lys	Met	Ala	Ser 100	Pro	Val	Ser	Ile	Thr 105	Trp	Ala	Val	Arg	Pro 110	Leu	Thr
30	Leu	Ser	Ser 115	Arg	Суѕ	Val	Thr	Ala 120	Gly	Thr	Ser	Cys	Ser 125	Phe	Pro	Ala
35	Gly	Ala 130	Ala	Arg	Pro	Asp	Pro 135	Ser	Tyr	Ala	Cys	Leu 140	Thr	Pro	Суз	Asp
	Ala 145	Pro	Thr	Ser	Pro	Ser 150	Leu	Ser	Thr	Arg	Ser 155	Val	Arg	Thr	Pro	Thr 160
40	Pro	Ala	Thr	Ser	Gln 165	Thr	Pro	Trp	Cys	Val 170	Pro	Ala	Cys	Arg	Lys 175	Gly
• .	Ala	Arg	Thr	Pro 180	Ala	Arg	Val	Thr	Pro 185	Gly	Ala	Leu	Trp	Ser 190	Val	Thr
45	Ser	Leu	Phe 195	Lys	Ala	Leu	Ser	Pro 200	Gly	Ala		Ile			Arg	Ser
50	Pro	Glu 210	.Ser	Leu	Val	Ser	Thr 215	Arg	Lys	Ser	Ala	Asn 220	Met	Trp	Thr	Gly
	Ser 225	Arg	Arg	Arg												
55	(2)	INFY	ORMAT	TON	FOR	SEO	TD N	ກຸ້າ	43-							
							11	· ,	- .							

(i) SEQUENCE CHARACTERISTICS:

60

(A) LENGTH: 74 amino acids

(B) TYPE: amino acid

```
(D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 743:
      Cys Lys Leu His Ser Gln Pro Trp Gln Ala Ala Leu Phe Glu Lys Thr
                                          10
     Arg Leu Cys Gly Ala Thr Leu Ile Ala Pro Arg Trp Leu Leu Thr
                                      25
10
     Ala Ala His Cys Leu Lys Pro Arg Tyr Ile Val His Leu Gly 31- His
     Asn Leu Gln Lys Glu Glu Gly Cys Glu Gln Thr Arg Thr Ala Thr Glu
15
      Ser Phe Pro His Pro Gly Phe Asn Asn Ser
20
      (2) INFORMATION FOR SEQ ID NO: 744:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 81 amino acids
25
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 744:
     Val Leu Gln Gly Arg Tyr Phe Ser Pro Ile Leu Glu Met Arg Arg Leu
30
                                          10
     Arg Pro Glu Gly Xaa Xaa Asn Leu Pro Gly Gly Ser Arg Ala Glm Lys
                                      25
35
     Glu Pro Arg Gln Asp Leu Thr Leu Val Leu Trp Pro His Cys Pro His
     Phe Ala Met Thr Arg Ser Tyr Val Pro Thr Lys Gln Cys Met Val 31m
40
     Gly Ser Phe Tyr Cys Ile Phe Ile Phe Lys Gly Pro Val Glm Asm Trp
     Cys
45
      (2) INFORMATION FOR SEQ ID NO: 745:
50
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 211 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
55
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 745:
     Met Pro Ile Ile Asp Gln Val Asn Pro Glu Leu His Asp Phe Met Gln
                                         10
60
     Ser Ala Glu Val Gly Thr Ile Phe Ala Leu Ser Trp Leu Ile Thr Trp
```

			21)				25	,				30		
5	Phe Gi	.y ≝ 3	s 7a] 5	Leu	Ser	Asp	Phe 40	arg	r His	7al	Val	Arg 45		Тут	Asp
J	75e 75	s le :	u Als	Cys	#£s	7 2 20 55	Leu	Xet	Pro	Ile	T∕∓ 60		λla	Ala	(Va)
10	Ile Va 65	l Le	ı Tyı	: Arg	Glu 70	Gln	`Glu	/al	Leu	Asp 75		Asp	Cys	Asp	Met 80
	Ala Se	r Va	l His	: #1s 25	Leu	Leu	Ser	Sln	Ile 90		Gln	Asp	Leu	Pro 95	
15	9lu Th	r Le	: Ile 100	Ser	æş	Xaa	Slu	Thr 105	Phe	Leu	Phe	Ser	Phe	Pro	His
20	Pro As	n Len 11	: Leu	Glγ	Arg	3 ≖0	Leu 120	220	Asn	Ser	Lys	Leu 125	Arg	Gly	Arg
	3lm Pr 13	s Le: C	. Leu	Ser	Lys	The 135	Leu	Ser	طتق	His	Gln 140	Pro	Ser	Arg	Gly
25	Leu Il 145	e Tr	: Cys	C/s	Gly 150	Ser	Gly	Хаа	Arg	Gly 155	Leu	Leu	Arg	Pro	Glu 160
	Asp Ar	g Thi	Lys	Asp 165	Val	Leu	Thr	Ŀys	Pro 170	λzg	Thr	Asn	Arg	Phe 175	Val
30	Lys le	i Als	Val 180	Met	Gly	Leu	Thr	Val 135	Ala	Leu	Gly	Ala	Ala 190	Ala	Leu
35	Ala Va	l Val 195	Lys	Ser	Ala	Leu	Glu 200	شتي	Ala	?ro	Lys	Phe 205	Gln	Leu	Gln
	Deu Phi II:														
40	(2) 그림	FORMA	rici;	FCR	<u>ತಾ</u> ರೆ	נו כב	70: 7	45:							
45			. (A) LI B) Ti D) TX	ENGT: (PE: (POL)	i: 70 amir XGY:	o ami no ac lina	ino a cid ear	acid		746	i:			
50	Cys Pro	Glu	Phe	Phe 5	Ile	Pro	Ala	Thr	Leu 10	Pro	Cys	Pro	Phe	Val 15	Phe
	Ala Phe	e Thr	Ser 20	Glu .	Ala	Ser	Ser .	A≃g 25	٨la	Tyz	Leu	Thr	Gln . 30	Arg	Gly
55	Pro Gly	Gly 35	Leu	Ala (Gln .	Asn .	Leu 1 40	Met	Pro	Leu	Pro	Val (Gly	Phe	Trp
50	Met Gly	Ser	Leu	Pzo .	2 x o :	9ro '	Trp (Cys '	Trp .	Arg :	Lys 60	Trp '	Val:	Ser	Gl ų

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```
Ala Cys Ser Cys Phe Cys
      65
5
      (2) INFORMATION FOR SEQ ID NO: 747:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 20 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 747:
     Gly Phe Gly Ser Val Ser Ala Ala Gly Arg Arg Ser Gly Gly Thr Trp
15
                        5
                                           10
      Gln Pro Val Gln
20
      (2) INFORMATION FOR SEQ ID NO: 748:
             (i) SEQUENCE CHARACTERISTICS:
25
                     (A) LENGTH: 16 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 748:
      Pro Gly Gly Leu Ala Val Gly Ser Arg Trp Trp Ser Arg Ser Leu Thr
30
                        5
                                           10
35
      (2) INFORMATION FOR SEQ ID NO: 749:
40
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 30 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 749:
45
      Leu Glu Pro Ser Arg Gln Arg Arg Pro Arg Arg Arg Gly Gly Thr Ser
        1
      Arg Pro Glu Thr Asp Gln Arg Ala Lys Cys Trp Arg Gln Leu
50
                   20
                                        25
       (2) INFORMATION FOR SEQ ID NO: 750:
55
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 11 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
 60
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 750:
```

	Val C	ys L	eu A	urg C	ys G 5	ln A	lsn A	rg M	iet (Glu A 10	\sn					
5																
	(2) I	NFOF	TAM	ION F	OR S	EQ I	ED NC): 75	51:							
10				(B (D) LE) TY) TO	NGTH PE: POLO	: 36 amin GY:	7 am o ac line	ino id ar	acid	s NO:	751	:			
15	Met A	la i	Ala	Cys 1	Thr i	Ala /	Arg 2	Arg	Pro	Gly . 10	Arg (Gly (Gln I	Pro I	Leu 1	Val
	Val F	Pro '	Val	Ala i 20	Asp :	Xaa	Gly	Pro	Val 25	Ala	Lys i	Ala .	Ala 1	Leu (30	Cys	Ala
20	Ala 2	Kaa .	Ala 35	Gly .	Ala	Phe	Ser	Pro 40	Ala	Ser	Thr '	Thr	Thr '	Thr	Arg	Arg
25	His I	Leu 50	Ser	Ser	Arg	Asn	Arg 55	Pro	Glu	Gly	Lys '	Val 60	Leu	Glu	Thr	Val
	Gly \ 65	Val	Phe	Glu	Val	Pro 70	Lys	Gln	Asn	Gly	Lys 75	Tyr	Glu	Thr	Gly	Gln 80
30	Leu :	Phe	Leu	His	Ser 85	Ile	Phe	Gly	Tyr	Arg 90	Gly	Val	Val	Leu	Phe 95	Pro
25	Trp	Gln	Ala	Arg 100	Leu	Xaa	Asp	Arg	Asp 105	Val	Ala	Ser	Ala	Ala 110	Pro	Glu
35	Lys	Ala	Glu 115		Pro	Ala		Нis 120		Ser	Lys	Glu	Val 125	Lys	Gly	Lys
40	Thr	His 130	Thr	Tyr	Туг	Gln	Val 135	Leu	Ile	Asp	Ala	Arg 140	Asp	Суз	Pro	His
٠.	Ile 145	Ser	Gln	Arg	Ser	Gln 150		Glu	Ala	Val	Thr 155	Phe	Leu	Ala	Asn	His 160
45	Asp	Asp	Ser	Arg	Ala 165		Tyr	Ala	Ile	Pro 170	Gly	Leu	Asp	Tyr	Val 175	Ser
50	His	Glu	Asp	1le 180		Pro	туг	Thr	Ser 189		: Asp	Gln	Val	Pro 190	Ile	Gln
50	His	Glu	Leu 195		Glu	Arg	j Ph∈	Let 200		1 Тут	: Asp	Glr	Thr 205	Lys	Ala	Pro
55	Pro	Phe 210		l Ala	Arg	g Glu	1 Thi 215		ı Ar	g Ala	a Trp	Glr 220	n Glu	Lys	Asr	n His
	Pro 225		Le	u Glu	Le.	230		va.	l Hi	s Arg	g Glu 235		Thr	: Glu	AST	1le 240
				_		_		_		1			- 01.	1.	. 61,	n Aen

		245	250	255											
	Ser His Val Tyr 260		Tyr Cys Ile Arg Leu 265	Glu Asn Leu Asp 270											
5	Ser Asp Val Val 275	Gln Leu Arg	Glu Arg His Trp Arg 280	g Ile Phe Ser Leu 285											
0	Ser Gly Thr Let 290	Glu Thr Val 295	Arg Gly Arg Gly Va	l Val Gly Arg Glu)											
	Pro Val Leu Ser 305	: Lys Glu Glr 310	Pro Ala Phe Gln Ty 315	r Ser Ser His Val											
15	Ser Leu Gln Ala	a Ser Ser Gly 325	His Met Trp Gly Th	r Phe Arg Phe Glu 335											
30	Arg Pro Asp Gl		a Asp Val Arg Ile Pr 345	o Pro Phe Ser Leu 350											
20	Glu Ser Asn Ly 355	s Asp Glu Ly	s Thr Pro Pro Ser Gl 360	y Leu His Trp 365											
25	(2) INFORMATIO	N FOR SEQ ID	NO: 752:												
30	(2) INFORMATION FOR SEQ ID NO: 752: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 33 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 752: Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val														
35	Met Ala Ala C	ys Thr Ala Ai 5	rg Arg Pro Gly Arg G	ly Gln Pro Leu Val 15											
		la Asp Xaa G 20	ly Pro Val Ala Lys A 25	la Ala Leu Cys Ala 30											
40	Ala														
45	(2) INFORMATI	ON FOR SEQ I	D NO: 753:												
50	•	(B) TYPE: (D) TOPOLO	: 33 amino acids amino acid	753:											
55	1	5	Arg Arg Pro Gly Arg 10	13											
<i>J</i> J	Val Pro Val	Ala Asp Xaa (20	Gly Pro Val Ala Lys 25	Ala Ala Leu Cys Ala 30											
60	Ala														

```
(2) INFORMATION FOR SEQ ID NO: 754:
  5
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 33 amino acids
                      (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 754:
 10
       Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val
       Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala
 15
       Ala
 20
        (2) INFORMATION FOR SEQ ID NO: 755:
               (i) SEQUENCE CHARACTERISTICS:
- 25
                      (A) LENGTH: 33 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 755:
  30
        Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val
                                             10
        Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala
                                         25
  35
                     20
        Ala
  40
        (2) INFORMATION FOR SEQ ID NO: 756:
                (i) SEQUENCE CHARACTERISTICS:
                       (A) LENGTH: 33 amino acids
  45
                       (B) TYPE: amino acid
                       (D) TOPOLOGY: linear
                (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 756:
        Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val
   50
                                              10
         Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala
                      20
   55
         Ala
```

	(2) INFORMATION FOR SEQ ID NO: 757:
5	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 35 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 757:
10	Val Leu Glu Thr Val Gly Val Phe Glu Val Pro Lys Gln Asn Gly Lys 1 5 10 15
	Tyr Glu Thr Gly Gln Leu Phe Leu His Ser Ile Phe Gly Tyr Arg Gly 20 25 30
15	Val Val Leu 35
20	(2) INFORMATION FOR SEQ ID NO: 758:
25	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 16 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 758:
30	Gly Leu Asp Tyr Val Ser His Glu Asp Ile Leu Pro Tyr Thr Ser Thr 1 5 10 15
35	(2) INFORMATION FOR SEQ ID NO: 759:
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 19 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 759:
45	Asp Val His Arg Glu Thr Thr Glu Asn Ile Arg Val Thr Val Ile Pro 1 5 10 15
	Phe Tyr Met
50	
	(2) INFORMATION FOR SEQ ID NO: 760:
55	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 21 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 760:
60	

```
1
                                          10
                                                              15
     Gln Leu Arg Glu Arg
                  20
5
     (2) INFORMATION FOR SEQ ID NO: 761:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 761:
15
     Pro Ala Phe Gln Tyr Ser Ser His Val Ser Leu Gln Ala Ser Ser Gly
                       5
                               10
     His Met Trp Gly Thr Phe Arg Phe Glu Arg
20
                  20
      (2) INFORMATION FOR SEQ ID NO: 762:
25
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 11 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 762:
      Ser Leu Cys Cys Pro Glu Gly Ala Glu Gly Cys
                       5
35
      (2) INFORMATION FOR SEQ ID NO: 763:
             (i) SEQUENCE CHARACTERISTICS:
40
                    (A) LENGTH: 12 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 763:
45
      Gln Leu Lys Lys Thr His Tyr Asp Arg Pro Cys Pro
50
      (2) INFORMATION FOR SEQ ID NO: 764:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 12 amino acids
                    (B) TYPE: amino acid
55
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 764:
      Gln Leu Lys Lys Thr His Tyr Asp Arg Pro Cys Pro
60
```

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734

	(2)	INFO	RMAT	NOI	FOR	SEQ	ID N	10: 7	65:							
5			(i) S	()	A) L B) T	CHAP ENGTI YPE: OPOLA	H: 1 ami	70 ar	nino cid		is					
			(xi)							EQ II	OM	: 76	5:			
10	Ala 1	Gln	Arg	Lys	Lys 5	Glu	Met	Val	Leu	Ser 10	Glu	Lys	Val	Ser	Gln 15	Leu
15	Met	Glu	Trp	Thr 20	Asn	Lys	Arg	Pro	Val 25	Ile	Arg	Met	Asn	Gly 30	Asp	Lys
	Phe	Arg	Arg 35	Leu	Val	Lys	Ala	Pro 40	Pro	Arg	Asn	Tyr	Ser 45	Val	Ile	Val
20	Met	Phe 50	Thr	Ala	Leu	Gln	Leu 55	His	Arg	Gln	Cys	Val 60	Val	Cys	Lys	Gln
25	Ala 65	_	Glu	Glu	Phe	Gln 70	Ile	Leu	Ala	Asn	Ser 75	Trp	Arg	Тут	Ser	Ser 80
	Ala	Phe	Thr	Asn	Arg 85		Phe	Phe	Ala	Met 90		Asp	Phe	Asp	Glu 95	Gly
30	Ser	Asp	Val	Phe 100		Met	Leu	Asn	Met 105		Ser	Ala	Pro	Thr 110	Phe	Ile
	Asn	Phe	Pro 115		Lys	Gly	Lys	Pro 120		Arg	Gly	Asp	Thr 125		Glu	Leu
35	Gln	Val	. Arg	Gly	Phe	e Ser	Ala 135	,	Gln	Ile	Ala	140		Ile	Ala	Ası
40	Arg 145		Asp	Val	. Asr	11e		Val	Ile	Arg	155) Asn	Met	Ala	160
	Arg	Tr	Arg	Phe	169		. Val	. Ser	Val	170						
45	(2)	IN	FORM	YTIO	1 FOE	R SEÇ) ID	NO:	766:	:						
			(i)	SEQ		Е СНІ										
50					(B) (D)	LENG TYPE TOPO	: am LOGY	ino : li	acid near							
						CE D										
55		t Va	l Va	l Ala		u Lei 5	ı Ile	e Va	l Cy:	s Asp 10		l Pro	o Sei	c Ala	a Ser	
	(2)) IN	FORM	ATIO	N FO	R SEX	Q ID	NO:	767	:						

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 16 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
 5
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 767:
      Ala Gln Arg Lys Lys Glu Met Val Leu Ser Glu Lys Val Ser Gln Leu
10
15
      (2) INFORMATION FOR SEQ ID NO: 768:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 17 amino acids
                     (B) TYPE: amino acid
20
                  · (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 768:
      Met Glu Trp Thr Asn Lys Arg Pro Val Ile Arg Met Asn Gly Asp Lys
                                           10
                                                               15
25
      Phe
30
       (2) INFORMATION FOR SEQ ID NO: 769:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 56 amino acids
35
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 769:
      Arg Arg Leu Val Lys Ala Pro Pro Arg Asn Tyr Ser Val Ile Val Met
 40
                                           10
       Phe Thr Ala Leu Gln Leu His Arg Gln Cys Val Val Cys Lys Gln Ala
                                        25
       Asp Glu Glu Phe Gln Ile Leu Ala Asn Ser Trp Arg Tyr Ser Ser Ala
 45
                                    40
       Phe Thr Asn Arg Ile Phe Phe Ala
            50
 50
       (2) INFORMATION FOR SEQ ID NO: 770:
 55
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 31 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 770:
 60
```

```
Met Val Asp Phe Asp Glu Gly Ser Asp Val Phe Gln Met Leu Asn Met
                       5
     Asn Ser Ala Pro Thr Phe Ile Asn Phe Pro Ala Lys Gly Lys Pro
                                      25
5
      (2) INFORMATION FOR SEQ ID NO: 771:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 37 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 771:
15
      Lys Arg Gly Asp Thr Tyr Glu Leu Gln Val Arg Gly Phe Ser Ala Glu
      Gln Ile Ala Arg Trp Ile Ala Asp Arg Thr Asp Val Asn Ile Arg Val
20
                                       25
      Ile Arg Pro Pro Asn
               35
25
      (2) INFORMATION FOR SEQ ID NO: 772:
              (i) SEQUENCE CHARACTERISTICS:
30
                     (A) LENGTH: 44 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 772:
35
      Tyr Ala Gly Pro Leu Met Leu Gly Leu Leu Leu Ala Val Ile Gly Gly
      Leu Val Tyr Leu Arg Arg Val Ile Trp Asn Phe Ser Leu Ile Lys Leu
40
      Asp Gly Leu Leu Gln Leu Cys Val Leu Cys Leu Leu
 45
       (2) INFORMATION FOR SEQ ID NO: 773:
              (i) SEQUENCE CHARACTERISTICS:
 50
                      (A) LENGTH: 17 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 773:
       Asp Ala Val Phe Lys Gly Phe Ser Asp Cys Leu Leu Lys Leu Gly Asp
 55
                                            10
       Ser
 60
```

	(2) INFORMATION FOR SEQ ID NO: 774:
5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids (B) TYPE: amino acid
10	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 774:
	Cys Gln Glu Gly Ala Lys Asp Met Trp Asp Lys Leu Arg Lys Glu Ser 1 5 10 15
15	Lys Asn Leu Asn 20
20	(2) INFORMATION FOR SEQ ID NO: 775: (i) SEQUENCE CHARACTERISTICS:
25	(A) LENGTH: 16 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 775: Val Leu Leu Val Ser Leu Ser Ala Ala Leu Ala Thr Trp Leu Ser Phe
30	1 5 10 15
35	(2) INFORMATION FOR SEQ ID NO: 776: (i) SEQUENCE CHARACTERISTICS:
40	(A) LENGTH: 48 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 776:
	Met Gly Leu Lys Leu Asn Gly Arg Tyr Ile Ser Leu Ile Leu Ala Val 1 5 10 15
45	Gln Ile Ala Tyr Leu Val Gln Ala Val Arg Ala Ala Gly Lys Cys Asp 20 25 30
50	Ala Val Phe Lys Gly Phe Ser Asp Cys Leu Leu Lys Leu Gly Asp Ser 35 40 45
55	
	(2) INFORMATION FOR SEQ ID NO: 777:
60	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 90 amino acids

		(xi)				GY:			EQ II	NO:	777	' :			
5	Pro Ala	a Ala	Trp i	Asp . 5	Asp	Lys	Thr	Asn	Ile 10	Lys	Thr	Val	Cys	Thr 15	Tyr
10	Trp Gl	ı Asp	Phe 1 20	His	Ser	Cys	Thr	Val 25	Thr	Ala	Leu	Thr	Asp 30	Суѕ	Gln
10	Glu Gl	y Ala 35	Lys :	Asp	Met	Trp	Asp 40	Lys	Leu	Arg	Lys	Glu 45	Ser	Lys	Asn
15	Leu As:		Gln	Gly	Ser	Leu 55	Phe	Glu	Leu	Cys	Gly 60	Ser	Gly	Asn	Gly
	Ala Al 65	a Gly	Ser	Leu	Le u 70	Pro	Ala	Phe	Pro	Val 75	Leu	Leu	Val	Ser	Leu 80
20	Ser Al	a Ala	Leu	Ala 85	Thr	Trp	Leu	Ser	Phe 90						
25	(2) IN		rion Seoue						i.e.						
30			() ()	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	43 a no a lin	mino cid ear	aci		: 77	8:			
35	Met Gl	y Leu	Lys	Leu 5	Asn	Gly	Arg	Tyr	Ile 10		Leu	Ile	Leu	Ala 15	Va]
	Gln Il	e Ala	Tyr 20	Leu	Val	Gln	Ala	Val 25		Ala	Ala	Gly	Lys 30		Asp
40	Ala Va	1 Phe 35		Gly	Phe	Ser	Asp 40		Leu	Leu	Lys	Leu 45		Asp	Sei
٠.	Xaa Xa	aa Xaa 50	Xaa	Xaa	Pro	Ala 55		Tr	Asp	Asp	Lys 60		. Asn	lle	Ly:
45	Thr Va	al Cys	Thr	Tyr	Trp 70		Asp	Phe	His	75		Thr	. Val	. Thr	* Ala 80
50	Leu T			85					90)				95	,
	Lys G	lu Ser	100		Leu	ı Asr	Ile	105		, Ser	Leu	ı Phe	9 Glu 110		ı Cy:
55	Gly S	er Gly 119		Gly	Ala	Ala	120		. Leu	ı Le	ı Pro	125		Pro	Va
	Leu L 1	eu Val 30	l Ser	Leu	Ser	: Ala 135		Let	ı Ala	Thi	140		u Sei	r Phe	•

	(2) INFORMATION FOR SEQ ID NO. 775.
5	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 34 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 779:
10	Met Asn Ser Ala Ala Gly Phe Ser His Leu Asp Arg Arg Glu Arg Val 1 5 10 15
15	Leu Lys Leu Gly Glu Ser Phe Glu Lys Gln Pro Arg Cys Ala Ser Thr 20 25 30
10	Leu Cys
20	(2) INFORMATION FOR SEQ ID NO: 780:
25	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 28 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 780:
30	Thr Ile Tyr Pro Thr Glu Glu Glu Leu Gln Ala Val Gln Lys Ile Val 1 5 10 15
	Ser Ile Thr Glu Arg Ala Leu Lys Leu Val Ser Asp 20 25
35	
	(2) INFORMATION FOR SEQ ID NO: 781:
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids (B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 781:
45	Arg Ala Leu Lys Gly Val Leu Arg Val Gly Val Leu Ala Lys Gly Leu 1 5 10 15
	Leu Leu Arg Gly Asp Arg Asn Val Asn Leu Val Leu Leu Cys
50	20 25 30
	(2) INFORMATION FOR SEQ ID NO: 782:
55	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 39 amino acids (B) TYPE: amino acid
	(D) TOPOLOGY: linear
60	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 782:

```
Ala Leu Ala Ala Leu Arg His Ala Lys Trp Phe Gln Ala Arg Ala Asn
                                         10
     Gly Leu Gln Ser Cys Val Ile Ile Ile Arg Ile Leu Arg Asp Leu Cys
5
                  20
     Gln Arg Val Pro Thr Trp Ser
              35
10
      (2) INFORMATION FOR SEQ ID NO: 783:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 17 amino acids
15
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 783:
      Gly Asp Ala Leu Arg Arg Val Phe Glu Cys Ile Ser Ser Gly Ile Ile
20
                                         10
                       5
        1
      Leu
25
      (2) INFORMATION FOR SEQ ID NO: 784:
              (i) SEQUENCE CHARACTERISTICS:
30
                    (A) LENGTH: 16 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 784:
35
      Leu Ala Phe Arg Gln Ile His Lys Val Leu Gly Met Asp Pro Leu Pro
                       5 . 10
 40
       (2) INFORMATION FOR SEQ ID NO: 785:
 45
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 342 amino acids
                    . (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 785:
 50
       Thr. Ile Tyr Pro Thr Glu Glu Glu Leu Gln Ala Val Gln Lys Ile Val
                                       10
                        5
       Ser Ile Thr Glu Arg Ala Leu Lys Leu Val Ser Asp Ser Leu Ser Glu
 55
                                       25
       His Glu Lys Asn Lys Asn Lys Glu Gly Asp Asp Lys Lys Glu Gly Gly
                35
 60
```

	Lys	Asp 50	Arg	Ala	Leu	Lys	Gly 55	Val	Leu	Arg	Val	Gly 60	Val	Leu	Ala	Lys
5	Gly 65	Leu	Leu	Leu	Arg	Gly 70	Asp	Arg	Asn	Val	Asn 75	Leu	Val	Leu	Leu	Cys 80
	Ser	Glu	Lys	Pro	Ser 85	Lys	Thr	Leu	Leu	Ser 90	Arg	Ile	Ala	Glu	Asn 95	Leu
10	Pro	Lys	Gln	Leu 100	Ala	Val	Ile	Ser	Pro 105	Glu	Lys	Tyr	Asp	Ile 110	Lys	Cys
15	Ala	Val	Ser 115	Glu	Ala	Ala	Ile	Ile 120	Leu	Asn	Ser	Cys	Val 125	Glu	Pro	Lys
	Met	Gln 130		Thr	Ile	Thr	Leu 135	Thr	Ser	Pro	Ile	Ile 140	Arg	Glu	Glu	Asn
20	Met 145	_	Glu	Gly	Asp	Val 150	Thr	Ser	Gly	Met	Val 155	Lys	Asp	Pro	Pro	Asp 160
	Val	Leu	Asp	Arg	Gln 165	Lys	Cys	Leu	Asp	Ala 170		Ala	Ala	Leu	Arg 175	His
25	Ala	Lys	Trp	Phe 180		Ala	Arg	Ala	Asn 185		Leu	Gln	Ser	Cys 190		Ile
30	Ile	e Ile	Arg 195	Ile	Leu	Arg	Asp	200		Gln	Arg	Val	Pro 205		Trp	Ser
J 0	Asp	210		Ser	Trp	Ala	Met 215		Leu	. Let	ı Val	Glu 220		Ala	Ile	Ser
35	Ser 225		a Ser	s Ser	Pro	Gln 230		Pro	Gly	/ Asp	235		Arg	j Arg	Val	Phe 240
	Glı	u Cy:	s Ile	e Ser	Ser 245		IĮ	ą Ile	e Lev	250		Ser	Pro	Gly	255	Leu
40	Ası	p Pr	o Cy:	s Glu 260		Asp	Pro	o Phe	Asp 265		r Leu	ı Ala	a Thi	270		: Asp
45	Gl	n Gl	n Ar		ı Ası	o Ile	e Th	r Sei 280		r Ala	a Gli) Phe	28!		ı Arç	g Leu
.5	Le	u Al 29		e Ar	g Gli	ı Ile	29		s Vai	l Le	u Gly	7 Mei 30		p Pro	Leu	ı Pro
50	30 30		t Se	r Gl	n Ar	31(n Il	e Hi	s As	n Ası 31		g Ly	s Ar	g Ar	arg 320
	As	pS∈	er As	p Gl	y Va 32		p G1	y Ph	e Gl	u Al 33		u Gl	y Ly	s Ly:	s Ası 33	p Lys 5
55	Ly	rs As	ар Ту	r As 34	_	n Ph	е									

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(i) SEQUENCE CHARACTERISTICS: .
                    (A) LENGTH: 24 amino acids
                    (B) TYPE: amino acid
5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 786:
     Met Gly Ser Gln His Ser Ala Ala Ala Arg Pro Ser Ser Cys Arg Arg
                                           10
                                                              15
10
     Lys Gln Glu Asp Asp Arg Asp Gly
                  20
15
      (2) INFORMATION FOR SEQ ID NO: 787:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 30 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 787:
      Leu Leu Ala Glu Arg Glu Glu Glu Ala Ile Ala Gln Phe Pro Tyr
25
                                           10
      Val Glu Phe Thr Gly Arg Asp Ser Ile Thr Cys Leu Thr Cys
                   20
                                       25
30
      (2) INFORMATION FOR SEQ ID NO: 788:
              (i) SEQUENCE CHARACTERISTICS:
35
                     (A) LENGTH: 34 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 788:
40
      Gln Gly Thr Gly Tyr Ile Pro Thr Glu Gln Val Asn Glu Leu Val Ala
                                           10
                        5 ·
        1
      Leu Ile Pro His Ser Asp Gln Arg Leu Arg Pro Gln Arg Thr Lys Gln
                                       25
                   20
45
      Tyr Val
 50
       (2) INFORMATION FOR SEQ ID NO: 789:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 55 amino acids
 55
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 789:
       Ala Arg Leu Asn Val Gly Arg Glu Ser Leu Lys Arg Glu Met Leu Lys
 60
                                           10
                         5
```

```
Ser Gln Gly Val Lys Val Ser Glu Ser Pro Met Gly Ala Arg His Ser
     Ser Trp Pro Glu Gly Ala Ala Phe Cys Lys Lys Val Gln Gly Ala Gln
                                  40
     Met Gln Phe Pro Pro Arg Arg
          50
10
     (2) INFORMATION FOR SEQ ID NO: 790:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 15 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 790:
20
     Ala Arg Leu Asn Val Gly Arg Glu Ser Leu Lys Arg Glu Met Leu
                                      10
                        5
25
      (2) INFORMATION FOR SEQ ID NO: 791:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 20 amino acids
30
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 791:
      Leu Lys Ser Gln Gly Val Lys Val Ser Glu Ser Pro Met Gly Ala Arg
35
                                         10
                        5
      His Ser Ser Trp
40
      (2) INFORMATION FOR SEQ ID NO: 792:
              (i) SEQUENCE CHARACTERISTICS:
45
                     (A) LENGTH: 17 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 792:
      Ala Phe Cys Lys Lys Val Gln Gly Ala Gln Met Gln Phe Pro Pro Arg
50
      Arg
55
       (2) INFORMATION FOR SEQ ID NO: 793:
 60
              (i) SEQUENCE CHARACTERISTICS:
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60

```
(A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 793:
5
     Ala Phe Cys Lys Lys Val Gln Gly Ala Gln Met Gln Phe Pro Pro Arg
                                      10
                       5
       1
     Arg
10
      (2) INFORMATION FOR SEQ ID NO: 794:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 37 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NG: 794:
20
      Val Gln Val Leu Glu Gln Leu Thr Asn Asn Ala Val Ala Glu Ser Arg
                                          10
      Phe Asn Asp Ala Ala Tyr Tyr Tyr Trp Met Leu Ser Met Gln Cys Leu
25
                                       25
     . Asp Ile Ala Gln Asp
               35
 30
       (2) INFORMATION FOR SEQ ID NO: 795:
              (i) SEQUENCE CHARACTERISTICS:
 35
                     (A) LENGTH: 34 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 795:
 40
       Pro Ala Gln Lys Asp Thr Met Leu Gly Lys Phe Tyr His Pie Gln Arg
       Leu Ala Glu Leu Tyr His Gly Tyr His Ala Ile His Arg His Thr Glu .
                                                            30
                                       25
 45
                    20
       Asp Pro
  50
        (2) INFORMATION FOR SEQ ID NO: 796:
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 27 amino acids
  55
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 796:
        Leu Ala Lys Gln Ser Lys Ala Leu Gly Ala Tyr Arg Leu Ala Arg His
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10 15 1 Ala Tyr Asp Lys Leu Arg Gly Leu Tyr Ile Pro 20 5 (2) INFORMATION FOR SEQ ID NO: 797: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 36 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 797: 15 Ala Arg Phe Gln Lys Ser Ile Glu Leu Gly Thr Leu Thr Ile Arg Ala Lys Pro Phe His Asp Ser Glu Glu Leu Val Pro Leu Cys Tyr Arg Cys 20 25 30 Ser Thr Asn Asn 35 25 (2) INFORMATION FOR SEQ ID NO: 798: (i) SEQUENCE CHARACTERISTICS: 30 (A) LENGTH: 73 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 798: 35 Pro Leu Leu Asn Asn Leu Gly Asn Val Cys Ile Asn Cys Arg Gln Pro 5 Phe Ile Phe Ser Ala Ser Ser Tyr Asp Val Leu His Leu Val Glu Phe 25 40 Tyr Leu Glu Glu Gly Ile Thr Asp Glu Glu Ala Ile Ser Leu Ile Asp 40 Leu Glu Val Leu Arg Pro Lys Arg Asp Asp Arg Gln Leu Glu Ile Cys 45 55 Lys Gln Gln Leu Pro Asp Ser Cys Gly . 70 50 (2) INFORMATION FOR SEQ ID NO: 799: (i) SEQUENCE CHARACTERISTICS: 55 (A) LENGTH: 29 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 799: Met Pro Tyr Ala Gln Trp Leu Ala Glu Asn Asp Arg Phe Glu Glu Ala 60

	1	5		10			15
5	Gln Lys Ala	Phe His L	ys Ala Gl	y Arg Gln 25	Arg Glu	Ala	
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15							
	Phe Ser Val	His Arg F 5	ro Glu Ti	r Leu Phe 10		Ser Arg	Phe Leu 15
20	Leu His Sen	Leu Pro I 20	ys Asp Ti	er Pro Ser 25	Gly Ile	Ser Lys 30	Val Lys
	Ile Leu Phe 35						

A. The indications made below relate to the microorganism referred to in the description on page 161 . line N/A .					
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Colle	ection				
Address of depositary institution (including postal code and country 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	v)				
Date of deposit March 27, 1997	Accession Number 97979				
C. ADDITIONAL INDICATIONS (leave blank if not applicable	le) This information is continued on an additional sheet				
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession")					
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A. The indications made below relate to the microorganism referred to in the description on page 162 . line N/A .					
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Col	flection				
Address of depositary institution (including postal code and count	try)				
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America					
Date of deposit April 4, 1997	Accession Number 97974				
C. ADDITIONAL INDICATIONS (leave blank if not applicate	ble) This information is continued on an additional sheet				
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)					
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A. The indications made below relate to the microorganism referred to in the description on page 162 . line N/A .					
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Coll	lection				
Address of depositary institution (including postal code and country	(ער				
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	,				
Date of deposit May 29, 1997	Accession Number 209080				
C. ADDITIONAL INDICATIONS (leave blank if not applicab	This information is continued on an additional sheet				
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications. e.g., "Accession Number of Deposit")					
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A. The indications made below relate to the microorganism referred to in the description on page 164 , line N/A						
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet					
Name of depositary institution American Type Culture C	Collection					
Address of depositary institution (including postal code and code	unury)					
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America						
Date of deposit December 3, 1997	Accession Number 209511					
C. ADDITIONAL INDICATIONS (leave blank if not applied	icable) This information is continued on an additional sheet					
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)						
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession						
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A. The indications made below relate to the microorganism referred to in the description on page 167 . line N/A							
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet						
Name of depositary institution American Type Culture Colle	ection						
Address of depositary institution (including postal code and country	v)						
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	Manassas, Virginia 20110-2209						
Date of deposit April 4, 1997	Accession Number 97975						
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A. The indications made below relate to the microorganism referred to in the description on page 167 , line N/A .					
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture	e Collection				
Address of depositary institution (including postal code and c	country)				
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America					
Date of deposit May 29, 1997	Accession Number 209081				
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A. The indications made below relate to the microorganism referred to in the description on page 171 . line N/A .					
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Col	lection				
Address of depositary institution (including postal code and count	(ימ				
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America					
Date of deposit April 4, 1997	Accession Number 97976				
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A. The indications made below relate to the microorganism referre on page 172 , line N/A	·				
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Coll	lection				
Address of depositary institution (including postal code and country 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	7)				
Date of deposit April 4, 1997	Accession Number 97977				
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A. The indications made below relate to the microorganism referre on page 172 , line N/A	d to in the description				
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet				
Name of depositary institution American Type Culture Coll	ection				
Address of depositary institution (including postal code and country	(יע				
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America					
Date of deposit May 29, 1997	Accession Number 209082				
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A. The indications made below relate to the microorganism referred to in the description on page 176 , line N/A .		
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
Name of depositary institution American Type Culture Coll	lection	
Address of depositary institution (including postal code and country	7)	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit April 28, 1997	Accession Number 209007	
C. ADDITIONAL INDICATIONS (leave blank if not applicab	ble) This information is continued on an additional sheet	
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A. The indications made below relate to the microorganism referred to in the description on page 176 , line N/A			
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet			
Name of depositary institution American Type Culture Collection			
Address of depositary institution (including postal code and country 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	n)		
Date of deposit May 29, 1997	Accession Number 209083		
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A. The indications made below relate to the microorganism referred to in the description on page 179 . line N/A		
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Name of depositary institution American Type Culture Collection		
Address of depositary institution (including postal code and count	ת)	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
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Name of depositary institution American Type Culture Coll	lection	
Address of depositary institution (including postal code and country	(ער	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit May 29, 1997	Accession Number 209084	
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	This information is continued on an additional sheet	
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A. The indications made below relate to the microorganism referred to in the description on page 180 , line N/A			
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet			
Name of depositary institution American Type Culture Coll	lection		
Address of depositary institution (including postal code and country	7)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	·		
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A. The indications made below relate to the microorganism referred to in the description on page 180 , line N/A .		
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
Name of depositary institution American Type Culture Coll	ection	
Address of depositary institution (including postal code and country	(ע	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit May 29, 1997	Accession Number 209085	
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A. The indications made below relate to the microorganism referred to in the description on page 182 , line N/A .			
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet			
Name of depositary institution American Type Culture Collection			
Address of depositary institution (including postal code and country 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	y)		
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A. The indications made below relate to the microorganism referred to in the description on page 186 . line N/A .		
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet		
Name of depositary institution American Type Culture Col	llection	
Address of depositary institution (including postal code and count	(רצי)	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit April 28, 1997	Accession Number 209011	
C. ADDITIONAL INDICATIONS (leave blank if not applicab	ble) This information is continued on an additional sheet	
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A. The indications made below relate to the microorganism referred to in the description on page 174 , line N/A			
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet		
Name of depositary institution American Type Culture Coll	lection		
Address of depositary institution (including postal code and country	n)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America			
Date of deposit April 7, 1998	Accession Number 209746		
C. ADDITIONAL INDICATIONS (leave blank if not applicable	the) This information is continued on an additional sheet		
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)			
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WO 98/54963 PCT/US98/11422

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What Is Claimed Is:

- 1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:
- (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, having biological activity;
 - (f) a polynucleotide which is a variant of SEQ ID NO:X;
 - (g) a polynucleotide which is an allelic variant of SEQ ID NO:X;
 - (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
- (i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.
- 2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.
- 3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

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- 4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.
- 5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
- The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the Nterminus.
- 7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.
- 8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.
 - 9. A recombinant host cell produced by the method of claim 8.
 - 10. The recombinant host cell of claim 9 comprising vector sequences.
- 11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:
- (a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;
- (c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (f) a full length protein of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

- (g) a variant of SEQ ID NO:Y;
- (h) an allelic variant of SEQ ID NO:Y; or
- (i) a species homologue of the SEQ ID NO:Y.
- 12. The isolated polypeptide of claim 11, wherein the secreted form or the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.
- 13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.
- 14. A recombinant host cell that expresses the isolated polypeptide of claim 11.
 - 15. A method of making an isolated polypeptide comprising:
- (a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and
 - (b) recovering said polypeptide.
 - 16. The polypeptide produced by claim 15.
- 17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11 or the polynucleotide of claim 1.
- 18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
- (a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.
- 19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
- (a) determining the presence or amount of expression of the polypeptide of claim 11 in a biological sample; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

- 20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:
 - (a) contacting the polypeptide of claim 11 with a binding partner; and
- (b) determining whether the binding partner effects an activity of the polypeptide.
 - 21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.
- 22. A method of identifying an activity in a biological assay, wherein the method comprises:
 - (a) expressing SEQ ID NO:X in a cell;
 - (b) isolating the supernatant;
 - (c) detecting an activity in a biological assay; and
 - (d) identifying the protein in the supernatant having the activity.
 - 23. The product produced by the method of claim 22.

PATENT COOPERATION TREATY

PCT

DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT (PCT Article 17(2)(a) and Rule 39)

International application No.	Applicant's or agent's file reference	IMPORTANT DECLARAT	Date of mailing (day/month/year)	
PCTAUSSE/11422	PZ007PCT	IMI OKIANI DECLARAT	ON 1 4 OCT 1998	
International Patent Classification (IPC) or both national classification and IPC Please See Continuation Sheet. Applicant HUMAN GENOME SCIENCES, INC. This International Searching Authority hereby declares, according to Article 17(2)(a), that no international search report will be established on the international application for the reasons indicated below. 1. The subject matter of the international application relates to: a	••	International filing date (day/mon	h/year) (Earliest) Priority Date (day/month/year)	
Please See Continuation Sheet. Applicant HUMAN GENOME SCIENCES, INC. This International Searching Authority hereby declares, according to Article 17(2)(a), that no international search report will be established on the international application for the reasons indicated below. 1. The subject matter of the international application relates to: a. cientific theories. b. mathematical theories. c. plant varieties. d. animal varieties. e. assembilly biological processes for the production of plants and animals, other than microbiological processes and the products of such processes. f. schemes, rules or methods of performing purely mental acts. schemes, rules or methods of playing games. i. methods for treatment of the animal body by surgery or therapy. j. methods for treatment of the human body by surgery or therapy. k. diagnostic methods practiced on the human or animal body. I. mere presentations of information. m. computer programs for which this International Searching Authority is not equipped to search prior art. 2. The failure of the following parts of the international application to comply with prescribed requirements prevents a meaningful search from being carried out:				
This International Searching Authority hereby declares, according to Article 17(2)(a), that no international search report will be established on the international application for the reasons indicated below. 1.	International Patent Classification (IPC) or both national classification and IPC Please See Continuation Sheet.			
The subject matter of the international application relates to: a	· · ·			
2. The failure of the following parts of the international application to comply with prescribed requirements prevents a meaningful search from being carried out: the description the claims the drawings 3. The failure of the nucleotide and/or amino acid sequence listing to comply with the prescribed requirements prevents a meaningful search from being carried out: it does not comply with the prescribed standard it is not in the prescribed machine readable form 4. Further comments: Please See Continuation Sheet. Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Weshington, D.C. 20231 Authorized offices BRIAN R. STANTON	1. The subject matter of the international application relates to: a. scientific theories. b. mathematical theories. c. plant varieties. d. animal varieties. e. essentially biological processes for the production of plants and animals, other than microbiological processes and the products of such processes. f. schemes, rules or methods of doing business. g. schemes, rules or methods of performing purely mental acts. h. schemes, rules or methods of playing games. i. methods for treatment of the human body by surgery or therapy. j. methods for treatment of the animal body by surgery or therapy. k. diagnostic methods practiced on the human or animal body.			
The failure of the nucleotide and/or amino acid sequence listing to comply with the prescribed requirements prevents a meaningful search from being carried out: X	2. The failure of the following parts of the international application to comply with prescribed requirements prevents a			
it does not comply with the prescribed standard it is not in the prescribed machine readable form 4. Further comments: Please See Continuation Sheet. Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Authorized offices Authorized offices ERIAN R. STANTON	the description			
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Authorized offices Authorized offices BRIAN R. STANTON	meaningful search from being carried out: it does not comply with the prescribed standard			
Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 BRIAN R. STANTON				
Facsimile No. (703) 305-3230 Telephone No. (703) 308-0196	Commissioner of Patents and Trademarks Box PCT Commissioner of Patents and Trademarks District Fourth Commissioner of Patents and Trademarks			
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DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

International application No. PCT/US98/11422

The International Patent Classification (IPC) or National Classification and IPC are as	s listed below:		
IPC(6): A01N 37/18, 43/04; C12Q 1/00, 1/02, 1/68; C12N 5/00, 5/06, 15/00, 15/06, 15/09, 15/10, 15/11; G01N 33/53			
US CL.: 435, 4, 7.1, 69.1, 70.1, 71.1, 172.3, 243, 320.1, 325, 410; 514/2, 44; 530/350, 387.1			
4. Further Comments (Continued):			
Applicant has not responded to the invitation to pay additional fees mailed on 04 August 1998. Therefore, the search would be conducted on the first appearing invention whihe includes claims 1-10, 14, and 15 in so far as these claims are drawn to the first ten (10) appearing nucleotide sequences. However, no meaningful search could be carried out on these sequences because the CRF that was received for this case on 15 June 1998 was technically defective and could not be used to conduct a search of the prior art.			
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